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Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE

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**UTILITY  
PATENT APPLICATION  
TRANSMITTAL**

(Only for new nonprovisional applications under 37C.F.R. § 1.53(b))

Attorney Docket No.

PC10315AGPR

First Named Inventor or Application Identifier

Mark D. Fidock

Title

Enzyme PDE xiv

Express Mail Label No.

EJ248205689US

**APPLICATION ELEMENTS**

See MPEP chapter 600 concerning utility patent application contents.

ADDRESS TO:

Assistant Commissioner for Patents  
Box Patent Application  
Washington, DC 20231

1. ☒ \*Fee Transmittal Form (e.g., PTO/SB/17)  
(Submit an original, and a duplicate for fee processing)
2. ☒ Specification [Total Pages (preferred arrangement set forth below)
- Descriptive title of the Invention
  - Cross References to Related Applications
  - Statement Regarding Fed sponsored R&D
  - Reference in Microfiche Appendix
  - Background of the Invention
  - Brief Summary of the Invention
  - Brief Description of the Drawings (if filed)
  - Detailed Description
  - Claim(s)
  - Abstract of the Disclosure
3. ☒ Drawing(s) (35 U.S.C. 11.3) [Total sheets 

4. ☒ Oath or Declaration [Total pages 

a. ☒ Newly executed (original)

b. ☐ Copy from a prior application (37 CFR §1.63(d))  
(for continuation/divisional with Box 17 completed)  
[Note Box 5 below]

i. ☐ **DELETION OF INVENTOR(S)**  
Signed statement attached deleting inventor(s) named in the prior application, see 37 C.F.R. §§1.63(d)(2) and 1.33(b).

5. ☐ Incorporation By Reference (useable if Box 4b is checked)  
The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 4b, is considered to be part of the disclosure of the accompanying application and is hereby incorporated by reference therein.

6. ☐ Microfiche Computer Program (Appendix)
7. Nucleotide and/or Amino Acid Sequence Submission (if applicable, all necessary)
- a. ☐ Computer Readable Copy
- b. ☐ Paper Copy (identical to computer copy)
- c. ☐ Statement verifying identity of above copies

**ACCOMPANYING APPLICATION PARTS**

8. ☒ Assignment Papers (cover sheet & document(s))
9. ☐ 37 C.F.R. §3.73(b) Statement ☐ Power of Attorney  
(when there is an assignee)
10. ☐ English Translation Document (if applicable)
11. ☐ Information Disclosure Statement (IDS)/PTO-1449 ☐ Copies of IDS Citations
12. ☒ Preliminary Amendment
13. ☒ Return Receipt Postcard (MPEP 503)  
(Should be specifically itemized)
14. ☐ \*Small Entity ☐ Statement filed in prior application, Status still proper and desired (PTO/SB/09-12)
15. ☒ Certified Copy of Priority Document(s)  
(if foreign priority is claimed)
16. ☐ Other:

\*NOTE FOR ITEMS 1 & 14: IN ORDER TO BE ENTITLED TO PAY SMALL ENTITY FEES, A SMALL ENTITY STATEMENT IS REQUIRED (37 C.F.R. § 1.27), EXCEPT IF ONE FILED IN A PRIOR APPLICATION IS RELIED UPON (37 C.F.R. § 1.28).

17. If a **CONTINUING APPLICATION**, check appropriate box, and supply the requisite information below and in a preliminary amendment:
- ☐ Continuation ☐ Divisional ☐ Continuation-in-part (CIP) of prior application No: \_\_\_\_\_ / \_\_\_\_\_
- Prior application information: Examiner \_\_\_\_\_ Group/Art Unit: \_\_\_\_\_

**18. CORRESPONDENCE ADDRESS**

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Express Mail No. EJ248205689US

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

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APPLICANT: Mark D. Fidock : Examiner: To Be Assigned

SERIAL NO.: To Be Assigned :

FILED: Herewith : Art Unit: To Be Assigned

FOR: Enzyme PDE xiv :

-----  
Assistant Commissioner For Patents  
Washington, D.C. 20231

Sir:

PRELIMINARY AMENDMENT

In the matter of the application being filed herewith, please amend the application as follows:

In the Claims:

Please amend the claims as follows:

Please cancel claims 7, 13-15, and 18-25.

In claim 3, please delete "or claim 2".

In claim 6, please replace "any one of claims 3 to 5" with --claim 3--.

In claims 8 and 9, please replace "any one of claims 3 to 7" with --claim 3--.

In claim 10, please replace the entirety of the claim with

--An assay method for identifying an agent that can affect PDE\_XIV activity, the assay method comprising:  
    contacting the agent with a PDE\_XIV protein having an amino acid sequence according to claim 1; and  
    measuring the activity of the PDE\_XIV protein;  
wherein a difference between PDE activity in the absence of the agent and PDE activity in the presence of the agent is indicative that the agent can affect PDE\_XIV activity.--

In claim 12, please delete "or claim 11" and "or expression".

In claims 16 and 17, after each occurrence of "PDE", please insert --\_XIV--.

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Please add the following claims 26-36.

26. An assay method for identifying an agent that can affect PDE\_XIV expression, the assay method comprising  
contacting the agent with a nucleotide sequence according to claim 3;  
and  
measuring the expression of PDE\_XIV;  
wherein a difference between PDE expression in the absence of the agent and PDE expression in the presence of the agent is indicative that the agent can affect PDE\_XIV expression.

27. An assay method according to claim 26 wherein the assay is to screen for agents useful in the treatment of disorders found in any one or more of the putamen, the Caudate nucleus of the brain, the Occipital lobe of the brain, the heart, ovary, the pituitary gland, kidney, liver, small intestine, thymus, skeletal muscle, leukocyte regions, dorsal root ganglia, uterus, cochlea, small intestine (duodenum), astrocytoma, and appendix.

28. A process comprising the steps of:  
(a) performing the assay according to claim 26;  
(b) identifying one or more agents that do affect PDE\_XIV expression; and  
(c) preparing a quantity of those one or more identified agents.

29. An isolated PDE\_XIV protein, wherein said protein shares at least 75% homology with SEQ ID NO:1, SEQ ID NO:3, or SEQ ID NO:5.

30. A PDE\_XIV protein of claim 29, wherein said protein shares at least 85% homology with SEQ ID NO:1, SEQ ID NO:3, or SEQ ID NO:5.

31. A PDE\_XIV protein of claim 29, wherein said protein shares at least 95% homology with SEQ ID NO:1, SEQ ID NO:3, or SEQ ID NO:5.

32. An isolated nucleic acid encoding a PDE\_XIV protein, wherein said nucleic acid shares at least 75% homology with SEQ ID NO:2, SEQ ID NO:4, or SEQ ID NO:6.

33. A nucleic acid of claim 32, wherein said nucleic acid shares at least 85% homology with SEQ ID NO:2, SEQ ID NO:4, or SEQ ID NO:6.

34. A nucleic acid of claim 32, wherein said nucleic acid shares at least 95% homology with SEQ ID NO:2, SEQ ID NO:4, or SEQ ID NO:6.

35. An isolated nucleic acid encoding a PDE\_XIV protein, wherein said nucleic acid hybridizes under high stringent conditions to a nucleic acid as set forth in SEQ ID NO:2, SEQ ID NO:4, or SEQ ID NO:6, or hybridizes to the antisense strand thereof.

36. An isolated PDE\_XIV protein encoded by a nucleic acid of claim 35.

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In the specification:

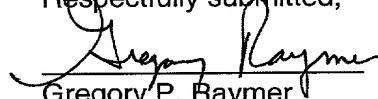
Please enter the following sentence as the first paragraph in the specification, immediately following the title:

-- This application is filed claiming priority from co-pending Great Britain Application No. 9828603.2, filed December 23, 1998, and Great Britain Application No. 9922123.6, filed September 17, 1999.--

A Notice of Allowance is courteously solicited.

Date: 12/22/99

Respectfully submitted,

  
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ENZYMES PDE xiv

## BACKGROUND OF THE PRESENT INVENTION

5 The present invention relates to an enzyme. The present invention also relates to a nucleotide sequence encoding same.

In particular, the present invention relates to novel nucleic acid sequences encoding novel phosphodiesterase enzymes.

10

The present invention also relates to the use of the novel nucleic acid and amino acid sequences in the diagnosis and treatment of disease.

15

The present invention also relates to the use of the novel nucleic acid and amino acid sequences to evaluate and/or to screen for agents that can modulate phosphodiesterase activity.

20

The present invention further relates to genetically engineered host cells that comprise or express the novel nucleic acid and amino acid sequences to evaluate and/or to screen for agents that can modulate phosphodiesterase activity.

## BACKGROUND ART

25

Cyclic nucleotides, such as cAMP and cGMP, are important intracellular second messengers. Cyclic nucleotide phosphodiesterases - otherwise known as PDEs - are a family of enzymes that catalyse the degradation of cyclic nucleotides and, in doing so, are one of the cellular components that regulate the concentration of cyclic nucleotides.

30

In recent years, at least ten PDE enzymes, as well as many subtypes of these enzymes, have been defined based on substrate affinity and cofactor requirements (Beavo JA and Reifsnyder DH, Trends Pharmacol. Sci. 11:150 [1990]; Beavo J, In: Cyclic Nucleotide Phosphodiesterases: Structure, Regulation and Drug Action., Beavo J and Housley MD (Eds.). Wiley:Chichester, pp. 3-15 [1990]).

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Examples of sub-families include: PDE1 (sometimes written as PDEI), Ca<sup>2+</sup>/calmodulin dependent; PDE2 (sometimes written as PDEII), cGMP-stimulated; PDE3 (sometimes written as PDEIII), cGMP-inhibited cAMP-hydrolysing; PDE4 (sometimes written as PDEIV), cAMP-specific, rolipram sensitive; PDE5 (sometimes written as PDEV), cGMP-specific; PDE6 (sometimes written as PDEVI), photoreceptor cGMP-specific; PDE7 (sometimes written as PDEVII), cAMP-specific, rolipram insensitive; PDE8 (sometimes written as PDEVIII), cAMP-specific IBMX insensitive; PDE9 (sometimes written as PDEIX), cGMP-specific IBMX insensitive; PDE10 (sometimes written as PDEX) dual substrate, IBMX sensitive.

PDEs contain a conserved C-terminal catalytic domain of ~270 amino acids (Charbonneau *et al* 1986 PNAS **83**(24):9308-12) and an N-terminal domain involved in regulating catalytic activity by binding cofactors, and in specifying subcellular localisation (Juilfs *et al* (1999) Rev Physiol Biochem Pharmacol **135**:67-104).

Each PDE family may contain two or more isoforms (i.e. there may be two or more PDE isoenzymes). By way of example, mammalian PDE IV, the homologue of the *Drosophila* Dunce gene (Chen CN *et al.*, Proc. Nat. Acad. Sci. (USA) 83:9313 [1986]), is known to have four isoforms in the rat (Swinnen JV *et al.*, Proc. Nat. Acad. Sci. (USA) 86:5325 [1989]). Human PDEs are also known to occur as isoforms and have splice variants. For example, the cloning of one human isoform of PDEIV from monocytes was reported in 1990 (Livi GP *et al.*, Mol. Cell. Bio., 10:2678 [1990]). By way of further example, other workers have independently cloned three splice variants of PDEIV, which are now designated hPDEIV-B1, hPDEIV-B2, and hPDEIV-B3.

Teachings on cyclic nucleotide phosphodiesterases can also be found in US-A-5932423 and US-A-5932465.

Teachings on a further cyclic nucleotide phosphodiesterase - namely CN PCDE8 - can be found in WO-A-97/35989. According to WO-A-97/35989, CN PCDE8 has two isozymes - which were designated CN PCDE8A and CN PCDE8B. The term "isozyme" is sometimes referred to in the art as "isoform".

According to WO-A-97/35989, many inhibitors of different PDEs have been identified and some have undergone clinical evaluation. For example, PDEIII inhibitors are being developed as antithrombotic agents, as antihypertensive agents and as cardiotonic agents useful in the treatment of congestive heart failure. Rolipram, a PDEIV inhibitor, has been used in the treatment of depression and other inhibitors of PDEIV are undergoing evaluation as anti-inflammatory agents. Rolipram has also been shown to inhibit lipopolysaccharide (LPS) induced TNF-alpha which has been shown to enhance HIV-1 replication *in vitro*. Therefore, rolipram may inhibit HIV-1 replication (Angel *et al* 1995 AIDS 9:1137-44). Additionally, based on its ability to suppress the production of TNF alpha and beta and interferon gamma, rolipram has been shown to be effective in the treatment of encephalomyelitis, the experimental animal model for multiple sclerosis (Sommer *et al*, 1995 Nat Med 1:244-248) and may be effective in the treatment of tardive dyskinesia (Sasaki *et al*, 1995 Eur J Pharmacol 282:71-76).

According to WO-A-97/35989, there are also non-specific PDE inhibitors such as theophylline, used in the treatment of bronchial asthma and other respiratory diseases, and pentoxifylline, used in the treatment of intermittent claudication and diabetes-induced peripheral vascular disease. Theophylline is thought to act on airway smooth muscle function as well as in an anti-inflammatory or immunomodulatory capacity in the treatment of respiratory diseases (Banner *et al* 1995 Respir J 8:996-1000) where it is thought to act by inhibiting both CN PDE cAMP and cGMP hydrolysis (Banner *et al* 1995 Monaldi Arch Chest Dis 50:286-292). Pentoxifylline, also known to block TNF-alpha production, may inhibit HIV-1 replication (Angel *et al supra*). A list of CN PDE inhibitors is given in Beavo 1995 *supra*.

It has been suggested that selective inhibitors of PDEs, in addition to their isozymes and their subtypes, will lead to more effective therapy with fewer side effects. For example, see the teachings in the reviews of Wieshaar RE *et al*, (J. Med. Chem., 28:537 [1985]), Giembycz MA (Biochem. Pharm., 43:2041 [1992]) and Lowe JA and Cheng JB (Drugs of the Future, 17:799-807 [1992]).

Thus, for some applications it is desirable to have a selective inhibition of an individual type of PDE. Hence, the cloning and expression of a novel PDE would greatly aid the discovery of selective inhibitors.

## 5 SUMMARY ASPECTS OF THE PRESENT INVENTION

Aspects of the present invention are presented in the claims and in the following commentary.

10 In brief, some aspects of the present invention relate to:

1. Novel amino acids.
2. Novel nucleotide sequences.
3. Assays using said novel sequences.
- 15 4. Compounds/compositions identified by use of said assays.
5. Expression systems comprising or expressing said novel sequences.
6. Methods of treatment based on said novel sequences.
7. Pharmaceutical compositions based on said novel sequences.

20 Other aspects concerning the amino acid sequence of the present invention and/or the nucleotide sequence of the present invention include: a construct comprising or capable of expressing the sequences of the present invention; a vector comprising or capable of expressing the sequences of the present invention; a plasmid comprising or capable of expressing the sequences of present invention; a tissue comprising or capable of expressing  
25 the sequences of the present invention; an organ comprising or capable of expressing the sequences of the present invention; a transformed host comprising or capable of expressing the sequences of the present invention; a transformed organism comprising or capable of expressing the sequences of the present invention. The present invention also encompasses methods of expressing the same, such as expression in a micro-organism; including methods  
30 for transferring same.

For ease of reference, aspects of the present invention are now discussed under appropriate section headings. However, the teachings under each section are not necessarily limited to each particular section.

5 In the following commentary references to "nucleotide sequence of the present invention" and "amino acid sequence of the present invention" refer respectively to any one or more of the nucleotide sequences presented or discussed herein and to any one or more of the amino acid sequences presented or discussed herein. Also, and as used herein, "amino acid sequence" refers to peptide or protein sequences and may refer to portions thereof. In  
10 addition, the term "amino acid sequence of the present invention" is synonymous with the phrase "polypeptide sequence of the present invention". Also, the term "nucleotide sequence of the present invention" is synonymous with the phrase "polynucleotide sequence of the present invention".

15 DETAILED ASPECTS OF THE PRESENT INVENTION

According to a first aspect of the present invention there is provided an amino acid sequence comprising the sequence presented as Formula I or a variant, homologue, fragment or derivative thereof; wherein the amino acid sequence is capable of displaying PDE activity.

20

Formula I is presented as follows:

**Formula I**

An amino acid sequence comprising any one or more of peptide sequences or amino acids Z1 - Z26, any of which peptide sequence or amino acid Z1 - Z26 may be separated from another  
 5 of said peptide sequence or amino acid Z1 - Z26 by a suitable peptide sequence or amino acid residue;

wherein:

10 Z1 = MSCLMVERCGE  
 Z2 = LFE  
 Z3 = P  
 Z4 = Q  
 Z5 = KVCMLGD  
 15 Z6 = RLRGQTGV  
 Z7 = AERRGSYPFIDFRLN  
 Z8 = TT  
 Z9 = SGEIGTKKKVKRLLSFQRYFHASRLLRGIIPQAPLHLLDEDYLGQARHMLSKVG  
 Z10 = WDFDIFLFDRLTNGNSLVTLCHLFN  
 20 Z11 = HGLIHHFKLDMVTLHRFLVMVQEDYH  
 Z12 =  
 NPYHNAVHAADVTQAMHCYLKEPKLASFLTPLDIMLGLLAAAHDVDHDPGVNQPFLLIKTNH  
 HLANLYQNMSVLENHHWRSTIGMLRESRLLAHLPKEMT  
 Z13 = QDIEQQLGSLILATDINRQNEFLTRLKAHLHNKDLRLE  
 25 Z14 = QDRHFMLQIALKCADICNPCRIWEMSKQWSERVCEEFYRQG  
 Z15 = LEQKFELEISPLCNQQKDSIPSIQIGFM  
 Z16 = YIVEPLFREWA  
 Z17 = FTGNSTLSENML  
 Z18 = HLAHNKAQWKSLL  
 30 Z19 = QHR  
 Z20 = RGS  
 Z21 = D  
 Z22 = AG  
 Z23 = E  
 35 Z24 = EQ  
 Z25 = EG  
 Z26 = P

and

40

wherein the amino acid sequence is made up of more than 200 amino acid residues.

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For convenience, we now present a Table indicating the codes used for the amino acids.

AMINO ACID	THREE LETTER ABBREVIATION	ONE LETTER SYMBOL
Alanine	Ala	A
Arginine	Arg	R
Asparagine	Asn	N
Aspartic acid	Asp	D
Cysteine	Cys	C
Glutamine	Gln	Q
Glutamic acid	Glu	E
Glycine	Gly	G
Histidine	His	H
Isoleucine	Ile	I
Leucine	Leu	L
Lysine	Lys	K
Methionine	Met	M
Phenylalanine	Phe	F
Proline	Pro	P
Serine	Ser	S
Threonine	Thr	T
Tryptophan	Trp	W
Tyrosine	Tyr	Y
Valine	Val	V
Any residue	Xaa	X

Here, Formula I - which can be expressed as being SEQ ID No. 9 - is a generic formula.

- 5 Formula I has been constructed from sequence analysis of the specific novel sequences presented herein.

For convenience, the PDE of the present invention is sometimes referred to as PDE\_XIV.

- We also sometimes refer to this enzyme as being PDEXIV. If the enzyme is obtainable from  
10 human we use the suffix or prefix HS (or HM) or hs (or hm). If the enzyme is obtainable from mouse we use the suffix or prefix MM (or Mm or mm).

The PDE of the present invention is capable of catalysing the degradation of cAMP.

- 15 For SEQ ID No. 9 any one or more of the amino acids may be an analogue thereof.

The term "analogue" as used herein means a sequence having a sequence similar to that of Formula I but wherein non-detrimental (i.e. not detrimental to enzymatic activity) amino acid substitutions or deletions have been made.

- 5 Preferably, the PDE enzyme of the present invention has at least two divalent cation binding motifs within the C-terminal catalytic domain and/or a conserved cAMP dependent protein kinase phosphorylation motif.

- 10 Preferably, the PDE enzyme of the present invention has at least two divalent cation binding motifs within the C-terminal catalytic domain and a conserved cAMP dependent protein kinase phosphorylation motif.

Preferably, the amino acid sequence of SEQ ID No. 9 comprises at least 5 of Z1 to Z26.

- 15 Preferably, the amino acid sequence of SEQ ID No. 9 comprises at least 10 of Z1 to Z26.

Preferably, the amino acid sequence of SEQ ID No. 9 comprises at least 15 of Z1 to Z26.

- 20 Preferably, the amino acid sequence of SEQ ID No. 9 comprises at least 20 of Z1 to Z26.

Preferably, the amino acid sequence of SEQ ID No. 9 comprises at least 25 of Z1 to Z26.

Preferably, the amino acid sequence of SEQ ID No. 9 comprises at least each of Z1 to Z26.

- 25 Preferably the amino acid sequence is made up of more than about 230 amino acid residues.

Preferably the amino acid sequence is made up of more than about 250 amino acid residues.

Preferably the amino acid sequence is made up of more than about 260 amino acid residues.

30

Preferably the amino acid sequence is made up of at least 268 amino acid residues.



Preferably, the amino acid sequence of Formula I comprises at least one or more of Z7, Z9, Z10, Z11, Z12, Z13, Z14, Z15, Z16, Z17 and Z18 or analogues thereof.

Preferably, the amino acid sequence of Formula I comprises at least each of Z7, Z9, Z10, Z11, Z12, Z13, Z14, Z15, Z16, Z17 and Z18 or analogues thereof.

Preferably, the amino acid sequence of Formula I comprises at least each of Z1, Z5, Z6, Z7, Z9, Z10, Z11, Z12, Z13, Z14, Z15, Z16, Z17 and Z18 or analogues thereof.

A preferred example of an amino acid sequence comprising the sequence presented as Formula I is the amino acid sequence shown as Formula II or a variant, homologue, fragment or derivative thereof.

### Formula II

Z1-X1-Z2-X2-Z3-X3-Z4-X4-Z5-X5-Z6-X6-Z7-X7-Z8-X8-Z9-X9-Z10-X10-Z11-X11-Z12-Z13-X12-Z14-X13-Z15-X14-Z16-X15-Z17-X16-Z18-X17-Z19-X18-Z20-X19-Z21-X20-Z22-X21-Z23-X22-Z24-X23-Z25-X24-Z26

wherein

each of Z1 - Z26 is as defined above; and

each of X1 - X24 is independently selected from an optional suitable peptide sequence or amino acid.

A more preferred example of an amino acid sequence comprising the sequence presented as Formula I is the amino acid sequence shown as Formula III or a variant, homologue, fragment or derivative thereof.

**Formula III**

5

Z1-X1-Z2-X2-Z3-X3-Z4-X4-Z5-X5-Z6-X6-Z7-X7-Z8-X8-Z9-X9-Z10-  
 X10-Z11-X11-Z12-Z13-X12-Z14-X13-Z15-X14-Z16-X15-Z17-X16-Z18-  
 X17-Z19-X18-Z20-X19-Z21-X20-Z22-X21-Z23-X22-Z24-X23-Z25-X24-  
 Z26

10

wherein

each of Z1 - Z26 is as defined above; and

- 15 X1 = V or I  
 X2 = S or N  
 X3 = E or D  
 X4 = a peptide comprising at least two or more of S, V, N  
 or A
- 20 X5 = V or I  
 X6 = P or R  
 X7 = N or S  
 X8 = H or Y  
 X9 = T or M
- 25 X10 = S or T  
 X11 = a peptide comprising at least two or more of G, H, S,  
 Q  
 X12 = a peptide comprising at least two or more of D, A, N,  
 V
- 30 X13 = E or D  
 X14 = S or T  
 X15 = H or R  
 X16 = G or S  
 X17 = a peptide comprising at least two or more of P, R, S,  
 35 N
- X18 = S or R  
 X19 = a peptide comprising at least two or more of S, G, P,  
 D, H, Q  
 X20 = H or L
- 40 X21 = a peptide comprising at least two or more of Q, G, T,  
 P, A  
 X22 = a peptide comprising at least two or more of S, E, T,  
 L  
 X23 = optional T
- 45 X24 = a peptide comprising at least two or more of D, S, A,  
 T

A more preferred example of an amino acid sequence comprising the sequence presented as Formula I is the amino acid sequence shown as Formula IV or a variant, homologue, fragment or derivative thereof.

5

**Formula IV**

Z1-X1-Z2-X2-Z3-X3-Z4-X4-Z5-X5-Z6-X6-Z7-X7-Z8-X8-Z9-X9-Z10-  
 X10-Z11-X11-Z12-Z13-X12-Z14-X13-Z15-X14-Z16-X15-Z17-X16-Z18-  
 X17-Z19-X18-Z20-X19-Z21-X20-Z22-X21-Z23-X22-Z24-X23-Z25-X24-

10 Z26

wherein

each of Z1 - Z26 is as defined above; and

15

X1 = V or I

X2 = S or N

X3 = E or D

X4 = SV or NA

20

X5 = V or I

X6 = P or R

X7 = N or S

X8 = H or Y

X9 = T or M

25

X10 = S or T

X11 = GH or SQ

X12 = DA or NV

X13 = E or D

X14 = S or T

30

X15 = H or R

X16 = G or S

X17 = PR or SN

X18 = S or R

X19 = SGSGPDH or GQ

35

X20 = H or L

X21 = QGT or PAP

X22 = SE or TL

X23 = optional T

X24 = DS or AT

40

An example of a variant of any one of the above Formulae is an amino acid sequence wherein the group ....HZ<sub>17</sub>GZ<sub>18</sub>PRZ<sub>19</sub>.... is substituted with ....RZ<sub>17</sub>SZ<sub>18</sub>SNZ<sub>19</sub>.... .

Preferred examples of an amino acid sequence according to the present invention which comprises the sequence presented as Formula I include the amino acids shown as: SEQ ID No. 1 or SEQ ID No. 3 or SEQ ID No. 5, or a variant, homologue, fragment or derivative of any thereof

5

It is to be noted that references to any one or more of the Formulae also apply equally to any one or more of: SEQ ID No. 1 or SEQ ID No. 3 or SEQ ID No. 5.

Preferably references to any one more of the Formulae herein mean any one or more of: SEQ ID No. 1 or SEQ ID No. 3 or SEQ ID No. 5.

10

More preferably references to any one or more of the Formulae mean a reference to SEQ ID No. 3 or 5.

According to a second aspect of the present invention there is provided an amino acid sequence comprising the sequence presented as Formula I.

15

According to a third aspect of the present invention there is provided a nucleotide sequence encoding the amino acid sequence of the present invention.

20

According to a fourth aspect of the present invention there is provided a nucleotide sequence comprising the sequence presented as SEQ ID No. 2 or SEQ ID No. 4 or SEQ ID No. 6, or a variant, homologue, fragment or derivative of any thereof.

According to a fifth aspect of the present invention there is provided a nucleotide sequence comprising the sequence presented as SEQ ID No. 2 or SEQ ID No. 4 or SEQ ID No. 6.

25

According to a sixth aspect of the present invention there is provided a nucleotide sequence that is capable of hybridising to the nucleotide sequence according to the present invention, or a sequence that is complementary thereto.

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According to a seventh aspect of the present invention there is provided a nucleotide sequence that is capable of hybridising to the nucleotide sequence according to the sixth aspect of the present invention, or a sequence that is complementary thereto.

- 5 According to an eighth aspect of the present invention there is provided a vector comprising the nucleotide sequence according to the present invention.

According to a ninth aspect of the present invention there is provided a host cell into which has been incorporated the nucleotide sequence according to the present invention.

10

- According to a tenth aspect of the present invention there is provided an assay method for identifying an agent that can affect PDE\_XIV activity or expression thereof, the assay method comprising contacting an agent with an amino acid according to the present invention or a nucleotide sequence according to the present invention; and measuring the activity or  
15 expression of PDE\_XIV; wherein a difference between a) PDE activity or expression in the absence of the agent and b) PDE activity or expression in the presence of the agent is indicative that the agent can affect PDE\_XIV activity or expression.

- Preferably the assay is to screen for agents useful in the treatment of disorders found in any  
20 one or more of the putamen, the Caudate nucleus of the brain, the Occipital lobe of the brain, the heart, ovary, the pituitary gland, kidney, liver, small intestine, thymus, skeletal muscle, leukocyte regions, dorsal root ganglia, uterus, cochlea, small intestine (duodenum), astrocytoma, and appendix.

- 25 According to an eleventh aspect of the present invention there is provided a process comprising the steps of: (a) performing the assay according to the present invention; (b) identifying one or more agents that do affect PDE\_XIV activity or expression; and (c) preparing a quantity of those one or more identified agents.

- 30 According to a twelfth aspect of the present invention there is provided a method of affecting *in vivo* PDE\_XIV activity or expression with an agent; wherein the agent is capable of affecting PDE\_XIV activity or expression in an *in vitro* assay method; wherein the *in vitro* assay method is the assay method of the present invention.

According to a thirteenth aspect of the present invention there is provided the use of an agent in the preparation of a pharmaceutical composition for the treatment of a disease or condition associated with PDE\_XIV, the agent is capable of having an effect on the activity or expression of PDE when assayed *in vitro* by the assay method of the present invention.

According to a fourteenth aspect of the present invention there is provided an enzyme capable of having an immunological reaction with an antibody raised against PDE\_XIV.

10 According to a fifteenth aspect of the present invention there is provided a nucleotide sequence coding for a PDE, wherein the nucleotide sequence is obtainable from NCIMB 40995 or NCIMB 40996 or NCIMB 41027.

15 According to a sixteenth aspect of the present invention there is provided a PDE wherein the PDE is expressable from a nucleotide sequence obtainable from NCIMB 40995 or NCIMB 40996 or NCIMB 41027.

20 According to a seventeenth aspect of the present invention there is provided the use of an agent which has an effect on the activity of PDE\_XIV or the expression thereof in the preparation of a pharmaceutical composition for the treatment of a disease or condition associated with PDE\_XIV.

According to a further aspect of the present invention there is provided a nucleotide sequence selected from:

25

- (a) the nucleotide sequence presented as any one of SEQ ID No. 2, 4, or 6;
- (b) a nucleotide sequence that is a variant, homologue, derivative or fragment of the nucleotide sequence presented as any one of SEQ ID No. 2, 4, or 6;
- (c) a nucleotide sequence that is the complement of the nucleotide sequence set out in  
30 any one of SEQ ID No. 2, 4, or 6;
- (d) a nucleotide sequence that is the complement of a variant, homologue, derivative or fragment of the nucleotide sequence presented as any one of SEQ ID No. 2, 4, or 6;

(e) a nucleotide sequence that is capable of hybridising to the nucleotide sequence set out in any one of SEQ ID No. 2, 4, or 6;

(f) a nucleotide sequence that is capable of hybridising to a variant, homologue, derivative or fragment of the nucleotide sequence presented as any one of SEQ ID No. 2, 4, or 6;

(g) a nucleotide sequence that is the complement of a nucleotide sequence that is capable of hybridising to the nucleotide sequence set out in any one of SEQ ID No. 2, 4, or 6;

(h) a nucleotide sequence that is the complement of a nucleotide sequence that is capable of hybridising to a variant, homologue, derivative or fragment of the nucleotide sequence presented as any one of SEQ ID No. 2, 4, or 6;

(i) a nucleotide sequence that is capable of hybridising to the complement of the nucleotide sequence set out in any one of SEQ ID No. 2, 4, or 6;

(j) a nucleotide sequence that is capable of hybridising to the complement of a variant, homologue, derivative or fragment of the nucleotide sequence presented as any one of SEQ ID No. 2, 4, or 6;

(k) a nucleotide sequence which is degenerate as a result of the genetic code to the nucleotides defined in (a), (b), (c), (d), (e), (f), (g), (h), (i), or (j);

(l) a nucleotide sequence comprising any one of (a), (b), (c), (d), (e), (f), (g), (h), (i), (j) and/or (k).

Other aspects of the present invention are now presented below.

An isolated nucleotide sequence or an isolated protein sequence according to the present invention.

A substantially pure nucleotide sequence or a substantially pure protein sequence according to the present invention.

An assay method for identifying an agent that can affect the expression pattern of the nucleotide sequence of the present invention or the activity of the expression product thereof, the assay method comprising: exposing the nucleotide sequence of the present invention or the expression product ("EP") thereof with an agent; determining whether the

agent modulates (such as affects the expression pattern or activity) the nucleotide sequence of the present invention or the expression product thereof.

An agent identified by the assay method of the present invention.

5

An agent identified by the assay method of the present invention, which agent has hitherto been unknown to have a PDE modulation effect in accordance with the present invention.

10

A process comprising the steps of: (a) performing the assay of the present invention; (b) identifying one or more agents that affect the expression pattern of the nucleotide sequence of the present invention or the activity of the expression product thereof; (c) preparing a quantity of those one or more identified agents.

15

A process comprising the steps of: (a) performing the assay according to the present invention; (b) identifying one or more agents that affect the expression pattern of the nucleotide sequence of the present invention or the activity of the expression product thereof; (c) preparing a pharmaceutical composition comprising one or more identified agents.

20

A process comprising the steps of: (a) performing the assay according to the present invention; (b) identifying one or more agents that affect the expression pattern of the nucleotide sequence of the present invention or the activity of the expression product thereof; (c) modifying one or more identified agents to cause a different effect on the expression pattern of the nucleotide sequence of the present invention or the activity of the expression product thereof.

25

Use of an agent identified by an assay according to the present invention in the manufacture of a medicament which affects the expression pattern of the nucleotide sequence of the present invention or the activity of the expression product thereof.

30

A method of treating a target (which target can be a mammal, preferably a human), which method comprises delivering (such as administering or exposing) to the target an effective



amount of an agent capable of modulating the expression pattern of the nucleotide sequence of the present invention or the activity of the expression product thereof.

5 A method of treating a target (which target can be a mammal, preferably a human), which method comprises delivering (such as administering or exposing) to the target an effective amount of an agent identified by an assay according to the present invention.

10 A method of inducing an immunological response in a subject, the method comprising administering to the subject the nucleotide sequence of the present invention or the expression product thereof.

#### PDE\_XIV

15 As explained above, the present invention relates to a novel PDE enzyme - which we have called PDE\_XIV - and to a nucleotide sequence encoding same. The present invention also relates to the use of the novel nucleic acid and amino acid sequences in the diagnosis and treatment of disease. The present invention also relates to the use of the novel nucleic acid and amino acid sequences to evaluate and/or to screen for agents that can modulate phosphodiesterase activity. The present invention further relates to genetically engineered  
20 host cells that comprise or express the novel nucleic acid and amino acid sequences to evaluate and/or to screen for agents that can modulate phosphodiesterase activity.

25 As used herein, the term "PDE\_XIV" refers to a novel family of PDEs which, up until now, were uncharacterised.

By way of example, we have identified a human PDE\_XIV (sometimes referred to as HSPDE\_XIV). We have also identified a mouse sequence - which we have called PDE\_XIV (sometimes referred to as MMPDE\_XIV).

30 For convenience, and unless otherwise stated, reference to PDE\_XIV will include reference to HSPDE\_XIV and/or MMPDE\_XIV.

PDE\_XIV is believed to be present in, and obtainable from, a variety of sources.

By way of example, PDE\_XIV is found in any one or more of the putamen, the Caudate nucleus of the brain, the Occipital lobe of the brain, the heart, ovary, the pituitary gland, kidney, liver, small intestine, thymus, skeletal muscle, leukocyte regions, dorsal root ganglia, uterus, cochlea, small intestine (duodenum), astrocytoma, and appendix.

We also believe that PDE\_XIV is also present in a number of other sources - such as for example: rat, bovine, ovine, porcine, and equine.

Preferably, the present invention covers mammalian PDE\_XIV which includes but is not limited to any of the above sources.

More preferably, the present invention covers human PDE\_XIV.

The PDE\_XIV may be the same as the naturally occurring form - for this aspect, preferably the PDE\_XIV is the non-native amino acid sequence (i.e. it is not present in its natural environment) - or is a variant, homologue, fragment or derivative thereof. In addition, or in the alternative, the PDE\_XIV is isolated PDE\_XIV and/or purified PDE\_XIV. The PDE\_XIV can be obtainable from or produced by any suitable source, whether natural or not, or it may be synthetic, semi-synthetic or recombinant.

The PDE\_XIV coding sequence may be the same as the naturally occurring form - for this aspect, preferably the PDE\_XIV coding sequence is the non-native nucleotide sequence (i.e. it is not present in its natural environment) - or is a variant, homologue, fragment or derivative thereof. In addition, or in the alternative, the PDE\_XIV coding sequence is an isolated PDE\_XIV coding sequence and/or a purified PDE\_XIV coding sequence. The PDE\_XIV coding sequence can be obtainable from or produced by any suitable source, whether natural or not, or it may be synthetic, semi-synthetic or recombinant.

PDE\_XIV and/or its coding sequence and/or a sequence capable of hybridising thereto is/are useful for testing the selectivity of drug candidates between different PDEs.

PDE\_XIV is believed to be able to catalyse the conversion of cAMP to AMP.

Preferred aspects of the present invention include a recombinant PDE\_XIV enzyme and a recombinant nucleotide sequence encoding a PDE\_XIV enzyme.

- 5 Preferably the recombinant PDE\_XIV enzyme and/or the recombinant nucleotide sequence of the present invention are a recombinant mammalian PDE\_XIV enzyme and/or a recombinant mammalian nucleotide sequence.

- 10 Preferably the recombinant PDE\_XIV enzyme and/or the recombinant nucleotide sequence of the present invention are a recombinant human PDE\_XIV enzyme and/or a recombinant human nucleotide sequence.

- 15 In accordance with the present invention, the recombinant PDE\_XIV enzyme has at least the formula presented as Formula I.

- 20 Either or both of the nucleotide sequence coding for PDE\_XIV or the enzyme PDE\_XIV itself may be used to screen for agents that can affect PDE\_XIV activity. In particular, the nucleotide sequence coding for PDE\_XIV or PDE\_XIV itself may be used to screen for agents that can inhibit PDE\_XIV activity. In addition, the nucleotide sequence coding for PDE\_XIV or the enzyme PDE\_XIV itself may be used to screen for agents that selectively affect PDE\_XIV activity, such as selectively inhibit PDE\_XIV activity.

- 25 Furthermore, the nucleotide sequence coding for PDE\_XIV or a sequence that is complementary thereto may also be used in assays to detect the presence of PDE\_XIV coding sequences in human cells. These assays would provide information regarding the tissue distribution of this enzyme and its biological relevance with respect to particular disease states.

- 30 The present invention also covers antibodies to PDE\_XIV (including a derivative, fragment, homologue or variant thereof). The antibodies for PDE\_XIV may be used in assays to detect the presence of PDE\_XIV in human cells. These assays would provide information regarding the tissue distribution of this enzyme and its biological relevance with respect to particular disease states.

In particular, any one or more of PDE\_XIV isozymes, the nucleotide sequences coding for same, the nucleotide sequences that are complementary to same, and the antibodies directed to same may be used in assays to screen for agents that selectively affect one of the isozymes.

- 5 These assays would provide information regarding the tissue distribution of each of the isozymes and to provide information regarding the biological relevance of each of the isozymes with respect to particular disease states. These assays would also allow workers to test for and identify agents that are useful to affect the expression of or activity of PDE\_XIV - such as in a particular tissue or in a particular disease state.

10

#### POLYPEPTIDE OF THE PRESENT INVENTION

- The term "polypeptide" - which is interchangeable with the term "protein" - includes single-chain polypeptide molecules as well as multiple-polypeptide complexes where individual  
15 constituent polypeptides are linked by covalent or non-covalent means.

Preferably, the polypeptide of the present invention is a single-chain polypeptide.

- Polypeptides of the present invention may be in a substantially isolated form. It will be  
20 understood that the polypeptide may be mixed with carriers or diluents which will not interfere with the intended purpose of the polypeptide and still be regarded as substantially isolated. A polypeptide of the present invention may also be in a substantially purified form, in which case it will generally comprise the polypeptide in a preparation in which more than 90%, e.g. 95%, 98% or 99% of the polypeptide in the preparation is a  
25 polypeptide of the present invention. Polypeptides of the present invention may be modified for example by the addition of histidine residues to assist their purification or by the addition of a signal sequence to promote their secretion from a cell as discussed below.

- Polypeptides of the present invention may be produced by synthetic means (e.g. as  
30 described by Geysen *et al.*, 1996) or recombinantly, as described below.

In a preferred embodiment, the amino acid sequence *per se* the present invention does not cover the native PDE\_XIV according to the present invention when it is in its natural

environment and when it has been expressed by its native nucleotide coding sequence which is also in its natural environment and when that nucleotide sequence is under the control of its native promoter which is also in its natural environment. For ease of reference, we have called this preferred embodiment the "non-native amino acid sequence".

5

- The terms "variant", "homologue" or "fragment" in relation to the amino acid sequence for the enzyme of the present invention include any substitution of, variation of, modification of, replacement of, deletion of or addition of one (or more) amino acid from or to the sequence providing the resultant enzyme has PDE\_XIV activity, preferably being at least as biologically active as the enzyme shown in the attached sequence listings. In particular, the term "homologue" covers homology with respect to structure and/or function. With respect to sequence homology, preferably there is at least 75%, more preferably at least 85%, more preferably at least 90% homology to the sequence shown as Formula I. More preferably there is at least 95%, more preferably at least 98%, homology to the sequence shown as Formula I.
- 10 More preferably there is at least 75%, more preferably at least 85%, more preferably at least 90% homology to any one of the sequences shown in the attached sequence listings. More preferably there is at least 95%, more preferably at least 98%, homology to any one of the sequence shown as shown in the attached sequence listings.
- 20 Typically, the types of amino acid substitutions that could be made should maintain the hydrophobicity/hydrophilicity of the amino acid sequence. Amino acid substitutions may be made, for example from 1, 2 or 3 to 10, 20 or 30 substitutions provided that the modified sequence retains the ability to act as a PDE enzyme in accordance with present invention. Amino acid substitutions may include the use of non-naturally occurring
- 25 analogues, for example to increase blood plasma half-life.

The amino acid sequence of the present invention may be produced by expression of a nucleotide sequence coding for same in a suitable expression system.

- 30 In addition, or in the alternative, the protein itself could be produced using chemical methods to synthesize a PDE amino acid sequence, in whole or in part. For example, peptides can be synthesized by solid phase techniques, cleaved from the resin, and purified by preparative high performance liquid chromatography (e.g., Creighton (1983) Proteins

Structures And Molecular Principles, WH Freeman and Co, New York NY). The composition of the synthetic peptides may be confirmed by amino acid analysis or sequencing (e.g., the Edman degradation procedure).

- 5 Direct peptide synthesis can be performed using various solid-phase techniques (Roberge JY *et al* (1995) Science 269: 202-204) and automated synthesis may be achieved, for example, using the ABI 431 A Peptide Synthesizer (Perkin Elmer) in accordance with the instructions provided by the manufacturer. Additionally, the amino acid sequence of PDE, or any part thereof, may be altered during direct synthesis and/or combined using chemical  
10 methods with a sequence from other subunits, or any part thereof, to produce a variant polypeptide.

In another embodiment of the invention, a PDE natural, modified or recombinant sequence may be ligated to a heterologous sequence to encode a fusion protein. For example, for  
15 screening of peptide libraries for inhibitors of PDE activity, it may be useful to encode a chimeric PDE protein expressing a heterologous epitope that is recognised by a commercially available antibody. A fusion protein may also be engineered to contain a cleavage site located between a PDE sequence and the heterologous protein sequence, so that the PDE may be cleaved and purified away from the heterologous moiety.

20 PDE may also be expressed as a recombinant protein with one or more additional polypeptide domains added to facilitate protein purification. Such purification facilitating domains include, but are not limited to, metal chelating peptides such as histidine-tryptophan modules that allow purification on immobilised metals (Porath J (1992) Protein  
25 Expr Purif 3 -.26328 1), protein A domains that allow purification on immobilised immunoglobulin, and the domain utilised in the FLAGS extension/affinity purification system (Immunex Corp, Seattle, WA). The inclusion of a cleavable linker sequence such as Factor XA or enterokinase (Invitrogen, San Diego, CA) between the purification domain and PDE is useful to facilitate purification. In a preferred aspect, the enzyme is expressed  
30 with a FLAG epitope tag in order to facilitate isolation and purification of the expressed enzyme.

Specific amino acid sequences of PDE\_XIV are shown as SEQ ID No. 1 and SEQ ID No. 3

and SEQ ID No. 5. However, the present invention encompasses amino acid sequences encoding other members from the PDE\_XIV family which would include amino acid sequences having at least 60% identity (more preferably at least 75% identity) to any one of the amino acid sequences. As indicated, suitable generic formulae for the PDE\_XIV family in accordance with the present invention are presented as Formula I.

Polypeptides of the present invention also include fragments of the presented amino acid sequence and variants thereof. Suitable fragments will be at least 5, e.g. at least 10, 12, 15 or 20 amino acids in size.

Polypeptides of the present invention may also be modified to contain one or more (e.g. at least 2, 3, 5, or 10) substitutions, deletions or insertions, including conserved substitutions. These aspects are discussed in a later section.

#### NUCLEOTIDE SEQUENCE OF THE PRESENT INVENTION

The term "nucleotide sequence" as used herein refers to an oligonucleotide sequence or polynucleotide sequence, and variants, homologues, fragments and derivatives thereof (such as portions thereof). The nucleotide sequence may be DNA or RNA which may be of genomic or synthetic or recombinant origin which may be double-stranded or single-stranded whether representing the sense or antisense strand.

Preferably, the term "nucleotide sequence" means DNA.

More preferably, the term "nucleotide sequence" means DNA prepared by use of recombinant DNA techniques (i.e. recombinant DNA).

In a preferred embodiment, the nucleotide sequence *per se* of the present invention does not cover the native nucleotide coding sequence according to the present invention in its natural environment when it is under the control of its native promoter which is also in its natural environment. For ease of reference, we have called this preferred embodiment the "non-native nucleotide sequence".

The nucleotide sequences of the present invention may include within them synthetic or modified nucleotides. A number of different types of modification to oligonucleotides are known in the art. These include methylphosphonate and phosphorothioate backbones, addition of acridine or polylysine chains at the 3' and/or 5' ends of the molecule. For the purposes of the present invention, it is to be understood that the nucleotide sequences described herein may be modified by any method available in the art. Such modifications may be carried out in to enhance the *in vivo* activity or life span of nucleotide sequences of the present invention.

10 The present invention also encompasses nucleotide sequences that are complementary to the sequences presented herein, or any derivative, fragment or derivative thereof. If the sequence is complementary to a fragment thereof then that sequence can be used a probe to identify similar coding sequences in other organisms etc.

15 The present invention also encompasses nucleotide sequences that are capable of hybridising to the sequences presented herein, or any derivative, fragment or derivative thereof.

The present invention also encompasses nucleotide sequences that are capable of hybridising to the sequences that are complementary to the sequences presented herein, or any derivative, fragment or derivative thereof.

20

The term "variant" also encompasses sequences that are complementary to sequences that are capable of hybridising to the nucleotide sequences presented herein.

25 Preferably, the term "variant" encompasses sequences that are complementary to sequences that are capable of hybridising under stringent conditions (e.g. 55°C and 0.2xSSC {1xSSC = 0.15 M NaCl, 0.015 Na<sub>3</sub> citrate pH 7.0}) to the nucleotide sequences presented herein.

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More preferably, the term "variant" encompasses sequences that are complementary to sequences that are capable of hybridising under high stringent conditions (e.g. 65°C and 0.1xSSC {1xSSC = 0.15 M NaCl, 0.015 Na<sub>3</sub> citrate pH 7.0}) to the nucleotide sequences presented herein.

5

The present invention also relates to nucleotide sequences that can hybridise to the nucleotide sequences of the present invention (including complementary sequences of those presented herein).

10 The present invention also relates to nucleotide sequences that are complementary to sequences that can hybridise to the nucleotide sequences of the present invention (including complementary sequences of those presented herein).

15 Also included within the scope of the present invention are polynucleotide sequences that are capable of hybridising to the nucleotide sequences presented herein under conditions of intermediate to maximal stringency.

20 In a preferred aspect, the present invention covers nucleotide sequences that can hybridise to the nucleotide sequence of the present invention, or the complement thereof, under stringent conditions (e.g. 55°C and 0.2xSSC).

In a more preferred aspect, the present invention covers nucleotide sequences that can hybridise to the nucleotide sequence of the present invention, or the complement thereof, under high stringent conditions (e.g. 65°C and 0.1xSSC).

25

Exemplary nucleic acids can alternatively be characterised as those nucleotide sequences which encode a PDE\_XIV protein and hybridise to any one or more of the DNA sequences shown in the attached sequence listings. Preferred are such sequences encoding PDE\_XIV which hybridise under high-stringency conditions to any one of the sequences shown in the  
30 attached sequence listings or the complement thereof.

Advantageously, the invention provides nucleic acid sequences which are capable of hybridising, under stringent conditions, to a fragment of any one of the sequences shown in

the attached sequence listings or the complement thereof. Preferably, the fragment is between 15 and 50 bases in length. Advantageously, it is about 25 bases in length.

The terms "variant", "homologue" or "fragment" in relation to the nucleotide sequence coding for the preferred enzyme of the present invention include any substitution of, variation of, modification of, replacement of, deletion of or addition of one (or more) nucleic acid from or to the sequence providing the resultant nucleotide sequence codes for or is capable of coding for an enzyme having PDE\_XIV activity, preferably being at least as biologically active as the enzyme encoded by any one of the sequences shown in the attached sequence listings. In particular, the term "homologue" covers homology with respect to structure and/or function providing the resultant nucleotide sequence codes for or is capable of coding for an enzyme having PDE\_XIV activity. With respect to sequence homology, preferably there is at least 75%, more preferably at least 85%, more preferably at least 90% homology to a nucleotide sequence coding for the amino acid sequence shown as Formula I. More preferably there is at least 95%, more preferably at least 98% homology to a nucleotide sequence coding for the amino acid sequence shown as Formula I. Preferably, with respect to sequence homology, preferably there is at least 75%, more preferably at least 85%, more preferably at least 90% homology to any one of the sequences shown in the attached sequence listings. More preferably there is at least 95%, more preferably at least 98%, homology to any one of the sequences shown in the attached sequence listings.

As indicated, the present invention relates to a DNA sequence (preferably a cDNA sequence) encoding PDE\_XIV. In particular, the present invention relates to cDNA sequences encoding PDE\_XIV.

The present invention also relates to DNA segments comprising the DNA sequence of any one of the sequences shown in the attached sequence listings or allelic variations of such sequences.

The present invention also relates to polypeptides produced by expression in a host cell into which has been incorporated the foregoing DNA sequences or allelic variations thereof.

The present invention also relates provides DNA comprising the DNA sequence of any one of the sequences shown in the attached sequence listings or an allelic variation thereof.

The present invention also relates to non-native DNA comprising the DNA sequence of any one of the sequences shown in the attached sequence listings or an allelic variation thereof.

A highly preferred aspect of the present invention relates to recombinant DNA comprising the DNA sequence of any one of the sequences shown in the attached sequence listings or an allelic variation thereof.

Polynucleotides of the present invention include nucleotide acid sequences encoding the polypeptides of the present invention. It will appreciated that a range of different polynucleotides encode a given amino acid sequence as a consequence of the degeneracy of the genetic code.

By knowledge of the amino acid sequences set out herein it is possible to devise partial and full-length nucleic acid sequences such as cDNA and/or genomic clones that encode the polypeptides of the present invention. For example, polynucleotides of the present invention may be obtained using degenerate PCR which will use primers designed to target sequences encoding the amino acid sequences presented herein. The primers will typically contain multiple degenerate positions. However, to minimise degeneracy, sequences will be chosen that encode regions of the amino acid sequences presented herein containing amino acids such as methionine which are coded for by only one triplet. In addition, sequences will be chosen to take into account codon usage in the organism whose nucleic acid is used as the template DNA for the PCR procedure. PCR will be used at stringency conditions lower than those used for cloning sequences with single sequence (non-degenerate) primers against known sequences.

Nucleic acid sequences obtained by PCR that encode polypeptide fragments of the present invention may then be used to obtain larger sequences using hybridisation library screening techniques. For example a PCR clone may be labelled with radioactive atoms and used to screen a cDNA or genomic library from other species, preferably other mammalian species. Hybridisation conditions will typically be conditions of medium to high stringency (for

example 0.03M sodium chloride and 0.03M sodium citrate at from about 50°C to about 60°C).

5 Degenerate nucleic acid probes encoding all or part of the amino acid sequence may also be used to probe cDNA and/or genomic libraries from other species, preferably other mammalian species. However, it is preferred to carry out PCR techniques initially to obtain a single sequence for use in further screening procedures.

10 In accordance with the present invention, PDE\_XIV polynucleotide sequences which encode PDE\_XIV, fragments of the polypeptide, fusion proteins or functional equivalents thereof, may be used to generate recombinant DNA molecules that direct the expression of PDE\_XIV in appropriate host cells. Due to the inherent degeneracy of the genetic code, other DNA sequences which encode substantially the same or a functionally equivalent amino acid sequence, may be used to clone and express PDE\_XIV. As will be understood  
15 by those of skill in the art, it may be advantageous to produce PDE-encoding nucleotide sequences possessing non-naturally occurring codons. Codons preferred by a particular prokaryotic or eukaryotic host (Murray E *et al* (1989) Nuc Acids Res 17:477-508) can be selected, for example, to increase the rate of PDE\_XIV expression or to produce recombinant RNA transcripts having desirable properties, such as a longer half-life, than  
20 transcripts produced from naturally occurring sequence.

Polynucleotide sequences of the present invention obtained using the techniques described above may be used to obtain further homologous sequences and variants using the techniques described above. They may also be modified for use in expressing the  
25 polypeptides of the present invention in a variety of host cells systems, for example to optimise codon preferences for a particular host cell in which the polynucleotide sequences are being expressed. Other sequence changes may be desired in order to introduce restriction enzyme recognition sites, or to alter the property or function of the polypeptides encoded by the polynucleotides.

30

Altered PDE\_XIV polynucleotide sequences which may be used in accordance with the invention include deletions, insertions or substitutions of different nucleotide residues resulting in a polynucleotide that encodes the same or a functionally equivalent PDE. The

protein may also have deletions, insertions or substitutions of amino acid residues which produce a silent change and result in a functionally equivalent PDE. Deliberate amino acid substitutions may be made on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophilicity, and/or the amphipathic nature of the residues as long as the biological activity of PDE is retained. For example, negatively charged amino acids include aspartic acid and glutamic acid; positively charged amino acids include lysine and arginine; and amino acids with uncharged polar head groups having similar hydrophilicity values include leucine, isoleucine, valine, glycine, alanine, asparagine, glutamine, serine, threonine, phenylalanine, and tyrosine.

Included within the scope of the present invention are alleles of PDE. As used herein, an "allele" or "allelic sequence" is an alternative form of PDE. Alleles result from a mutation, i.e., a change in the nucleic acid sequence, and generally produce altered mRNAs or polypeptides whose structure or function may or may not be altered. Any given gene may have none, one or many allelic forms. Common mutational changes which give rise to alleles are generally ascribed to deletions, additions or substitutions of amino acids. Each of these types of changes may occur alone, or in combination with the others, one or more times in a given sequence.

The term "allele" also includes genetic polymorphisms, such as SNPs (single nucleotide polymorphisms).

The nucleotide sequences of the present invention may be engineered in order to alter a PDE coding sequence for a variety of reasons, including but not limited to, alterations which modify the cloning, processing and/or expression of the gene product. For example, mutations may be introduced using techniques which are well known in the art, e.g., site-directed mutagenesis to insert new restriction sites, to alter glycosylation patterns or to change codon preference.

Polynucleotides of the present invention may be used to produce a primer, e.g. a PCR primer, a primer for an alternative amplification reaction, a probe e.g. labelled with a revealing label by conventional means using radioactive or non-radioactive labels, or the polynucleotides may be cloned into vectors. Such primers, probes and other fragments will

be at least 15, preferably at least 20, for example at least 25, 30 or 40 nucleotides in length, and are also encompassed by the term polynucleotides of the present invention as used herein.

5 Polynucleotides or primers of the present invention may carry a revealing label. Suitable labels include radioisotopes such as  $^{32}\text{P}$  or  $^{35}\text{S}$ , enzyme labels, or other protein labels such as biotin. Such labels may be added to polynucleotides or primers of the present invention and may be detected using by techniques known *per se*.

10 Polynucleotides such as a DNA polynucleotide and primers according to the present invention may be produced recombinantly, synthetically, or by any means available to those of skill in the art. They may also be cloned by standard techniques.

In general, primers will be produced by synthetic means, involving a step wise manufacture  
15 of the desired nucleic acid sequence one nucleotide at a time. Techniques for accomplishing this using automated techniques are readily available in the art.

Longer polynucleotides will generally be produced using recombinant means, for example using a PCR (polymerase chain reaction) cloning techniques. This will involve making a  
20 pair of primers (e.g. of about 15-30 nucleotides) to a region of the nucleotide sequence which it is desired to clone, bringing the primers into contact with mRNA or cDNA obtained from a fungal, plant or prokaryotic cell, performing a polymerase chain reaction under conditions which bring about amplification of the desired region, isolating the amplified fragment (e.g. by purifying the reaction mixture on an agarose gel) and  
25 recovering the amplified DNA. The primers may be designed to contain suitable restriction enzyme recognition sites so that the amplified DNA can be cloned into a suitable cloning vector.

DNA molecules may be modified to increase intracellular stability and half-life. Possible  
30 modifications include, but are not limited to, the addition of flanking sequences of the 5' and/or 3' ends of the molecule or the use of phosphorothioate or 2' O-methyl rather than phosphodiesterase linkages within the backbone of the molecule.

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As mentioned earlier, the present invention also relates to nucleotide sequences that are capable of hybridising to all or part of any one of the sequences shown in the attached sequence listings or an allelic variation thereof. These nucleotide sequences may be used in anti-sense techniques to modify PDE\_XIV expression. Alternatively, these sequences (or portions thereof) can be used as a probe, or for amplifying all or part of such sequence when used as a polymerase chain reaction primer.

In addition to the recombinant DNA sequences, genomic sequences are also of utility in the context of drug discovery. It may be valuable to inhibit the mRNA transcription of a particular isoform rather than to inhibit its translated protein. This may be true with PDE\_XIV, if there are splice variants and wherein those different splice variants may be transcribed from different promoters. There is precedent for multiple promoters directing the transcription of a mouse brain 2',3'-cyclic-nucleotide 3' phosphodiesterase (Kurihara T *et al.*, Biochem. Biophys. Res. Comm. 170:1074 [1990]).

Another utility of the invention is that the DNA sequences, once known, give the information needed to design assays to specifically detect isoenzymes or splice variants. Isozyme-specific PCR primer pairs are but one example of an assay that depends completely on the knowledge of the specific DNA sequence of the isozyme or splice variant. Such an assay allows detection of mRNA for the isozyme to access the tissue distribution and biological relevance of each isozyme to a particular disease state. It also allows identification of cell lines that may naturally express only one isozyme - a discovery that might obviate the need to express recombinant genes. If specific PDE\_XIV isozymes are shown to associated with a particular disease state, the invention would be valuable in the design of diagnostic assays to detect the presence of isozyme mRNA.

An abnormal level of nucleotide sequences encoding a PDE\_XIV in a biological sample may reflect a chromosomal aberration, such as a nucleic acid deletion or mutation. Accordingly, nucleotide sequences encoding a PDE\_XIV provide the basis for probes which can be used diagnostically to detect chromosomal aberrations such as deletions, mutations or chromosomal translocations in the gene encoding PDE. PDE\_XIV gene expression may be altered in such disease states or there may be a chromosomal aberration present in the region of the gene encoding a PDE\_XIV.

In an alternative embodiment of the invention, the coding sequence of PDE could be synthesized, in whole or in part, using chemical methods well known in the art (see Caruthers MH *et al* (1980) Nuc Acids Res Symp Ser 215-23, Horn T *et al* (1980) Nuc  
5 Acids Res Symp Ser 225-232).

#### NATURALLY OCCURRING

As used herein "naturally occurring" refers to a PDE\_XIV with an amino acid sequence  
10 found in nature.

#### ISOLATED/PURIFIED

As used herein, the terms "isolated" and "purified" refer to molecules, either nucleic or  
15 amino acid sequences, that are removed from their natural environment and isolated or separated from at least one other component with which they are naturally associated.

#### BIOLOGICALLY ACTIVE

As used herein "biologically active" refers to a PDE\_XIV according to the present invention - such as a recombinant PDE\_XIV - having a similar structural function (but not necessarily to the same degree), and/or similar regulatory function (but not necessarily to the same degree), and/or similar biochemical function (but not necessarily to the same degree) and/or immunological activity (but not necessarily to the same degree) of the  
20 naturally occurring PDE\_XIV. Specifically, a PDE\_XIV of the present invention has the ability to hydrolyse a cyclic nucleotide, which is one of the characteristic activities of the PDE enzyme of the present invention.  
25

#### IMMUNOLOGICAL ACTIVITY

30

As used herein, "immunological activity" is defined as the capability of the natural, recombinant or synthetic PDE\_XIV or any oligopeptide thereof, to induce a specific immune response in appropriate animals or cells and to bind with specific antibodies.



## DERIVATIVE

5 The term "derivative" as used herein in relation to the amino acid sequence includes chemical modification of a PDE\_XIV. Illustrative of such modifications would be replacement of hydrogen by an alkyl, acyl, or amino group.

## DELETION

10 As used herein a "deletion" is defined as a change in either nucleotide or amino acid sequence in which one or more nucleotides or amino acid residues, respectively, are absent.

## INSERTION/ADDITION

15 As used herein an "insertion" or "addition" is a change in a nucleotide or amino acid sequence which has resulted in the addition of one or more nucleotides or amino acid residues, respectively, as compared to the naturally occurring PDE.

## SUBSTITUTION

20 As used herein "substitution" results from the replacement of one or more nucleotides or amino acids by different nucleotides or amino acids, respectively.

## HOMOLOGUE

25 The term "homologue" with respect to the nucleotide sequence of the present invention and the amino acid sequence of the present invention may be synonymous with allelic variations of the sequences.

30 In particular, the term "homology" as used herein may be equated with the term "identity". Here, sequence homology with respect to the nucleotide sequence of the present invention and the amino acid sequence of the present invention can be determined by a simple "eyeball" comparison (i.e. a strict comparison) of any one or more of the sequences with

another sequence to see if that other sequence has at least 75% identity to the sequence(s). Relative sequence homology (i.e. sequence identity) can also be determined by commercially available computer programs that can calculate % homology between two or more sequences. A typical example of such a computer program is CLUSTAL.

5

% homology may be calculated over contiguous sequences, i.e. one sequence is aligned with the other sequence and each amino acid in one sequence directly compared with the corresponding amino acid in the other sequence, one residue at a time. This is called an “ungapped” alignment. Typically, such ungapped alignments are performed only over a relatively short number of residues (for example less than 50 contiguous amino acids).

Although this is a very simple and consistent method, it fails to take into consideration that, for example, in an otherwise identical pair of sequences, one insertion or deletion will cause the following amino acid residues to be put out of alignment, thus potentially resulting in a large reduction in % homology when a global alignment is performed. Consequently, most sequence comparison methods are designed to produce optimal alignments that take into consideration possible insertions and deletions without penalising unduly the overall homology score. This is achieved by inserting “gaps” in the sequence alignment to try to maximise local homology.

20

However, these more complex methods assign “gap penalties” to each gap that occurs in the alignment so that, for the same number of identical amino acids, a sequence alignment with as few gaps as possible - reflecting higher relatedness between the two compared sequences - will achieve a higher score than one with many gaps. “Affine gap costs” are typically used that charge a relatively high cost for the existence of a gap and a smaller penalty for each subsequent residue in the gap. This is the most commonly used gap scoring system. High gap penalties will of course produce optimised alignments with fewer gaps. Most alignment programs allow the gap penalties to be modified. However, it is preferred to use the default values when using such software for sequence comparisons. For example when using the GCG Wisconsin Bestfit package (see below) the default gap penalty for amino acid sequences is -12 for a gap and -4 for each extension.

30

Calculation of maximum % homology therefore firstly requires the production of an optimal alignment, taking into consideration gap penalties. A suitable computer program for carrying out such an alignment is the GCG Wisconsin Bestfit package (University of Wisconsin, U.S.A.; Devereux *et al.*, 1984, Nucleic Acids Research 12:387). Examples of other software than can perform sequence comparisons include, but are not limited to, the BLAST package (see Ausubel *et al.*, 1999 *ibid* – Chapter 18), FASTA (Atschul *et al.*, 1990, J. Mol. Biol., 403-410) and the GENWORKS suite of comparison tools. Both BLAST and FASTA are available for off-line and on-line searching (see Ausubel *et al.*, 1999 *ibid*, pages 7-58 to 7-60). However, for some applications it is preferred to use the GCG Bestfit program.

Although the final % homology can be measured in terms of identity, in some cases, the alignment process itself is typically not based on an all-or-nothing pair comparison. Instead, a scaled similarity score matrix is generally used that assigns scores to each pairwise comparison based on chemical similarity or evolutionary distance. An example of such a matrix commonly used is the BLOSUM62 matrix - the default matrix for the BLAST suite of programs. GCG Wisconsin programs generally use either the public default values or a custom symbol comparison table if supplied (see user manual for further details). It is preferred to use the public default values for the GCG package, or in the case of other software, the default matrix, such as BLOSUM62.

Once the software has produced an optimal alignment, it is possible to calculate % homology, preferably % sequence identity. The software typically does this as part of the sequence comparison and generates a numerical result.

As indicated, for some applications, sequence homology (or identity) may be determined using any suitable homology algorithm, using for example default parameters.

Advantageously, the BLAST algorithm is employed, with parameters set to default values.

The BLAST algorithm is described in detail at

[http://www.ncbi.nih.gov/BLAST/blast\\_help.html](http://www.ncbi.nih.gov/BLAST/blast_help.html). The search parameters are defined as follows, and are advantageously set to the defined default parameters.

Advantageously, "substantial homology" when assessed by BLAST equates to sequences which match with an EXPECT value of at least about 7, preferably at least about 9 and most preferably 10 or more. The default threshold for EXPECT in BLAST searching is usually 10.

5

BLAST (Basic Local Alignment Search Tool) is the heuristic search algorithm employed by the programs blastp, blastn, blastx, tblastn, and tblastx; these programs ascribe significance to their findings using the statistical methods of Karlin and Altschul (see [http://www.ncbi.nih.gov/BLAST/blast\\_help.html](http://www.ncbi.nih.gov/BLAST/blast_help.html)) with a few enhancements. The BLAST programs were tailored for sequence similarity searching, for example to identify homologues to a query sequence. The programs are not generally useful for motif-style searching. For a discussion of basic issues in similarity searching of sequence databases, see Altschul et al (1994) Nature Genetics 6:119-129.

10

15 The five BLAST programs available at <http://www.ncbi.nlm.nih.gov> perform the following tasks:

**blastp** compares an amino acid query sequence against a protein sequence database;

20 **blastn** compares a nucleotide query sequence against a nucleotide sequence database;

**blastx** compares the six-frame conceptual translation products of a nucleotide query sequence (both strands) against a protein sequence database;

25 **tblastn** compares a protein query sequence against a nucleotide sequence database dynamically translated in all six reading frames (both strands).

**tblastx** compares the six-frame translations of a nucleotide query sequence against the six-frame translations of a nucleotide sequence database.

30

BLAST uses the following search parameters:

HISTOGRAM Display a histogram of scores for each search; default is yes. (See parameter H in the BLAST Manual).

5 DESCRIPTIONS Restricts the number of short descriptions of matching sequences reported to the number specified; default limit is 100 descriptions. (See parameter V in the manual page). See also EXPECT and CUTOFF.

10 ALIGNMENTS Restricts database sequences to the number specified for which high-scoring segment pairs (HSPs) are reported; the default limit is 50. If more database sequences than this happen to satisfy the statistical significance threshold for reporting (see EXPECT and CUTOFF below), only the matches ascribed the greatest statistical significance are reported. (See parameter B in the BLAST Manual).

15 EXPECT The statistical significance threshold for reporting matches against database sequences; the default value is 10, such that 10 matches are expected to be found merely by chance, according to the stochastic model of Karlin and Altschul (1990). If the statistical significance ascribed to a match is greater than the EXPECT threshold, the match will not be reported. Lower EXPECT thresholds are more stringent, leading to fewer chance matches being reported. Fractional values are acceptable. (See parameter E in the BLAST  
20 Manual).

CUTOFF Cutoff score for reporting high-scoring segment pairs. The default value is calculated from the EXPECT value (see above). HSPs are reported for a database sequence only if the statistical significance ascribed to them is at least as high as would be ascribed  
25 to a lone HSP having a score equal to the CUTOFF value. Higher CUTOFF values are more stringent, leading to fewer chance matches being reported. (See parameter S in the BLAST Manual). Typically, significance thresholds can be more intuitively managed using EXPECT.

30 MATRIX Specify an alternate scoring matrix for BLASTP, BLASTX, TBLASTN and TBLASTX. The default matrix is BLOSUM62 (Henikoff & Henikoff, 1992). The valid alternative choices include: PAM40, PAM120, PAM250 and IDENTITY. No alternate

scoring matrices are available for BLASTN; specifying the MATRIX directive in BLASTN requests returns an error response.

5 STRAND Restrict a TBLASTN search to just the top or bottom strand of the database sequences; or restrict a BLASTN, BLASTX or TBLASTX search to just reading frames on the top or bottom strand of the query sequence.

10 FILTER Mask off segments of the query sequence that have low compositional complexity, as determined by the SEG program of Wootton & Federhen (1993) Computers and Chemistry 17:149-163, or segments consisting of short-periodicity internal repeats, as determined by the XNU program of Claverie & States (1993) Computers and Chemistry 17:191-201, or, for BLASTN, by the DUST program of Tatusov and Lipman (see <http://www.ncbi.nlm.nih.gov>). Filtering can eliminate statistically significant but biologically uninteresting reports from the blast output (e.g., hits against common acidic-, 15 basic- or proline-rich regions), leaving the more biologically interesting regions of the query sequence available for specific matching against database sequences.

20 Low complexity sequence found by a filter program is substituted using the letter "N" in nucleotide sequence (e.g., "NNNNNNNNNNNNNN") and the letter "X" in protein sequences (e.g., "XXXXXXXXXX").

Filtering is only applied to the query sequence (or its translation products), not to database sequences. Default filtering is DUST for BLASTN, SEG for other programs.

25 It is not unusual for nothing at all to be masked by SEG, XNU, or both, when applied to sequences in SWISS-PROT, so filtering should not be expected to always yield an effect. Furthermore, in some cases, sequences are masked in their entirety, indicating that the statistical significance of any matches reported against the unfiltered query sequence should be suspect.

30

NCBI-gi Causes NCBI gi identifiers to be shown in the output, in addition to the accession and/or locus name.

Most preferably, sequence comparisons are conducted using the simple BLAST search algorithm provided at <http://www.ncbi.nlm.nih.gov/BLAST>.

- Should Gap Penalties be used when determining sequence identity, then preferably the following parameters are used:

FOR BLAST	
GAP OPEN	0
GAP EXTENSION	0

FOR CLUSTAL	DNA	PROTEIN	
WORD SIZE	2	1	K triple
GAP PENALTY	10	10	
GAP EXTENSION	0.1	0.1	

- Other computer program methods to determine identify and similarity between the two sequences include but are not limited to the GCG program package (Devereux et al 1984 Nucleic Acids Research 12: 387 and FASTA (Atschul et al 1990 J Molec Biol 403-410).

#### POLYPEPTIDE VARIANTS AND DERIVATIVES

- The terms "variant" or "derivative" in relation to the amino acid sequences of the present invention includes any substitution of, variation of, modification of, replacement of, deletion of or addition of one (or more) amino acids from or to the sequence providing the resultant amino acid sequence has PDE activity, preferably having at least the same activity as any one of the polypeptides presented in the sequence listings.

The sequences of the present invention may be modified for use in the present invention. Typically, modifications are made that maintain the PDE activity of the sequence. Amino acid substitutions may be made, for example from 1, 2 or 3 to 10, 20 or 30 substitutions provided that the modified sequence retains the PDE activity. Amino acid substitutions

may include the use of non-naturally occurring analogues, for example to increase blood plasma half-life of a therapeutically administered polypeptide.

Conservative substitutions may be made, for example according to the Table below.

- 5 Amino acids in the same block in the second column and preferably in the same line in the third column may be substituted for each other:

ALIPHATIC	Non-polar	G A P
		I L V
	Polar - uncharged	C S T M
		N Q
	Polar - charged	D E
		K R
AROMATIC		H F W Y

- As indicated above, proteins of the invention are typically made by recombinant means, for example as described herein, and/or by using synthetic means using techniques well known to skilled persons such as solid phase synthesis. Variants and derivatives of such sequences include fusion proteins, wherein the fusion proteins comprise at least the amino acid sequence of the present invention being linked (directly or indirectly) to another amino acid sequence. These other amino acid sequences - which are sometimes referred to as fusion protein partners - will typically impart a favourable functionality - such as to aid extraction and purification of the amino acid sequence of the present invention. Examples of fusion protein partners include glutathione-S-transferase (GST), 6xHis, GAL4 (DNA binding and/or transcriptional activation domains) and  $\beta$ -galactosidase. It may also be convenient to include a proteolytic cleavage site between the fusion protein partner and the protein sequence of the present invention so as to allow removal of the latter. Preferably the fusion protein partner will not hinder the function of the protein of the present invention.



## POLYNUCLEOTIDE VARIANTS AND DERIVATIVES

The terms "variant" or "derivative" in relation to the nucleotide sequence of the present invention include any substitution of, variation of, modification of, replacement of, deletion  
5 of or addition of one (or more) nucleic acid from or to the sequence providing the resultant nucleotide sequence codes for a polypeptide having PDE activity, preferably having at least the same activity as sequences presented in the sequence listings.

As indicated above, with respect to sequence homology, preferably there is at least 75%,  
10 more preferably at least 85%, more preferably at least 90% homology to the sequences shown in the sequence listing herein. More preferably there is at least 95%, more preferably at least 98%, homology. Nucleotide homology comparisons may be conducted as described above. For some applications, a preferred sequence comparison program is the GCG Wisconsin Bestfit program described above. The default scoring matrix has a match value of 10 for each  
15 identical nucleotide and -9 for each mismatch. The default gap creation penalty is -50 and the default gap extension penalty is -3 for each nucleotide.

As used herein, the terms "variant", "homologue", "fragment" and "derivative" embrace allelic variations of the sequences.

20 The term "variant" also encompasses sequences that are complementary to sequences that are capable of hybridising to the nucleotide sequences presented herein.

## HYBRIDISATION

25 The term "hybridisation" as used herein shall include "the process by which a strand of nucleic acid joins with a complementary strand through base pairing" (Coombs J (1994) Dictionary of Biotechnology, Stockton Press, New York NY) as well as the process of amplification as carried out in polymerase chain reaction technologies as described in  
30 Dieffenbach CW and GS Dveksler (1995, PCR Primer, a Laboratory Manual, Cold Spring Harbor Press, Plainview NY).

Hybridisation conditions are based on the melting temperature ( $T_m$ ) of the nucleic acid binding complex, as taught in Berger and Kimmel (1987, Guide to Molecular Cloning Techniques, Methods in Enzymology, Vol 152, Academic Press, San Diego CA), and confer a defined "stringency" as explained below.

5

Stringency of hybridisation refers to conditions under which polynucleic acids hybrids are stable. Such conditions are evident to those of ordinary skill in the field. As known to those of skill in the art, the stability of hybrids is reflected in the melting temperature ( $T_m$ ) of the hybrid which decreases approximately 1 to 1.5°C with every 1% decrease in sequence homology. In general, the stability of a hybrid is a function of sodium ion concentration and temperature. Typically, the hybridisation reaction is performed under conditions of higher stringency, followed by washes of varying stringency.

As used herein, high stringency refers to conditions that permit hybridisation of only those nucleic acid sequences that form stable hybrids in 1 M Na<sup>+</sup> at 65-68 °C

Maximum stringency typically occurs at about  $T_m$ -5°C (5°C below the  $T_m$  of the probe).

High stringency at about 5°C to 10°C below the  $T_m$  of the probe. High stringency conditions can be provided, for example, by hybridisation in an aqueous solution containing 6x SSC, 5x Denhardt's, 1 % SDS (sodium dodecyl sulphate), 0.1 Na<sup>+</sup> pyrophosphate and 0.1 mg/ml denatured salmon sperm DNA as non specific competitor. Following hybridisation, high stringency washing may be done in several steps, with a final wash (about 30 min) at the hybridisation temperature in 0.2 - 0.1x SSC, 0.1 % SDS.

25

Moderate, or intermediate, stringency typically occurs at about 10°C to 20°C below the  $T_m$  of the probe.

Low stringency typically occurs at about 20°C to 25°C below the  $T_m$  of the probe.

30

As will be understood by those of skill in the art, a maximum stringency hybridisation can be used to identify or detect identical polynucleotide sequences while an intermediate (or

low) stringency hybridisation can be used to identify or detect similar or related polynucleotide sequences.

Moderate stringency refers to conditions equivalent to hybridisation in the above described solution but at about 60-62°C. In that case the final wash is performed at the hybridisation temperature in 1x SSC, 0.1 % SDS.

Low stringency refers to conditions equivalent to hybridisation in the above described solution at about 50-52°C. In that case, the final wash is performed at the hybridisation temperature in 2x SSC, 0.1 % SDS.

For some applications, it may be useful to use low to medium stringent conditions to identify similar enzymes coding sequences from different organisms.

It is understood that these conditions may be adapted and duplicated using a variety of buffers, e.g. formamide-based buffers, and temperatures. Denhardt's solution and SSC are well known to those of skill in the art as are other suitable hybridisation buffers (see, e.g. Sambrook, et al., eds. (1989) *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory Press, New York or Ausubel, et al., eds. (1990) *Current Protocols in Molecular Biology*, John Wiley & Sons, Inc.). Optimal hybridisation conditions have to be determined empirically, as the length and the GC content of the probe also play a role.

Polynucleotides of the invention capable of selectively hybridising to the nucleotide sequences presented herein, or to their complement, will be generally at least 70%, preferably at least 80 or 90% and more preferably at least 95% or 98% homologous to the corresponding nucleotide sequences presented herein over a region of at least 20, preferably at least 25 or 30, for instance at least 40, 60 or 100 or more contiguous nucleotides.

The term "selectively hybridisable" means that the polynucleotide used as a probe is used under conditions where a target polynucleotide of the invention is found to hybridize to the probe at a level significantly above background. The background hybridization may occur because of other polynucleotides present, for example, in the cDNA or genomic DNA library being screening. In this event, background implies a level of signal generated by interaction

between the probe and a non-specific DNA member of the library which is less than 10 fold, preferably less than 100 fold as intense as the specific interaction observed with the target DNA. The intensity of interaction may be measured, for example, by radiolabelling the probe, e.g. with  $^{32}\text{P}$ .

5

In a preferred aspect, the present invention covers nucleotide sequences that can hybridise to any one or more of the nucleotide sequences of the present invention under stringent conditions (e.g.  $60^{\circ}\text{C}$  and  $0.2\times\text{SSC}$  { $1\times\text{SSC} = 0.15\text{ M NaCl}, 0.015\text{ M Na}_3\text{ Citrate pH }7.0$ }).

10 In a more preferred aspect, the present invention covers nucleotide sequences that can hybridise to any one or more of the nucleotide sequences of the present invention under high stringent conditions (e.g.  $65^{\circ}\text{C}$  and  $0.1\times\text{SSC}$  { $1\times\text{SSC} = 0.15\text{ M NaCl}, 0.015\text{ M Na}_3\text{ Citrate pH }7.0$ }).

15 Where the polynucleotide of the present invention is double-stranded, both strands of the duplex, either individually or in combination, are encompassed by the present invention. Where the polynucleotide is single-stranded, it is to be understood that the complementary sequence of that polynucleotide is also included within the scope of the present invention.

20 Polynucleotides which are not 100% homologous to the sequences of the present invention but fall within the scope of the invention can be obtained in a number of ways. Other variants of the sequences described herein may be obtained for example by probing DNA libraries made from a range of individuals, for example individuals from different populations. In addition, other viral/bacterial, or cellular homologues particularly cellular homologues found  
25 in mammalian cells (e.g. rat, mouse, bovine and primate cells), may be obtained and such homologues and fragments thereof in general will be capable of selectively hybridising to the sequences shown in the sequence listing herein. Such sequences may be obtained by probing cDNA libraries made from or genomic DNA libraries from other animal species, and probing such libraries with probes comprising all or part of any one of the sequences in the attached  
30 sequence listings under conditions of medium to high stringency. Similar considerations apply to obtaining species homologues and allelic variants of the polypeptide or nucleotide sequences of the invention.

10 Variants and strain/species homologues may also be obtained using degenerate PCR which will use primers designed to target sequences within the variants and homologues encoding conserved amino acid sequences within the sequences of the present invention. Conserved sequences can be predicted, for example, by aligning the amino acid sequences from several  
5 variants/homologues. Sequence alignments can be performed using computer software known in the art. For example the GCG Wisconsin PileUp program is widely used.

10 The primers used in degenerate PCR will contain one or more degenerate positions and will be used at stringency conditions lower than those used for cloning sequences with single sequence primers against known sequences.

15 Alternatively, such polynucleotides may be obtained by site directed mutagenesis of characterised sequences. This may be useful where for example silent codon changes are required to sequences to optimise codon preferences for a particular host cell in which the polynucleotide sequences are being expressed. Other sequence changes may be desired in order to introduce restriction enzyme recognition sites, or to alter the property or function of the polypeptides encoded by the polynucleotides.

20 Polynucleotides of the invention may be used to produce a primer, e.g. a PCR primer, a primer for an alternative amplification reaction, a probe e.g. labelled with a revealing label by conventional means using radioactive or non-radioactive labels, or the polynucleotides may be cloned into vectors. Such primers, probes and other fragments will be at least 15, preferably at least 20, for example at least 25, 30 or 40 nucleotides in length, and are also encompassed by the term polynucleotides of the invention as used herein.

25 Polynucleotides such as a DNA polynucleotides and probes according to the invention may be produced recombinantly, synthetically, or by any means available to those of skill in the art. They may also be cloned by standard techniques.

30 In general, primers will be produced by synthetic means, involving a step wise manufacture of the desired nucleic acid sequence one nucleotide at a time. Techniques for accomplishing this using automated techniques are readily available in the art.

Longer polynucleotides will generally be produced using recombinant means, for example using a PCR (polymerase chain reaction) cloning techniques. This will involve making a pair of primers (e.g. of about 15 to 30 nucleotides) flanking a region of the lipid targeting sequence which it is desired to clone, bringing the primers into contact with mRNA or cDNA  
5 obtained from an animal or human cell, performing a polymerase chain reaction under conditions which bring about amplification of the desired region, isolating the amplified fragment (e.g. by purifying the reaction mixture on an agarose gel) and recovering the amplified DNA. The primers may be designed to contain suitable restriction enzyme recognition sites so that the amplified DNA can be cloned into a suitable cloning vector.

10

### REGULATORY SEQUENCES

Preferably, the polynucleotide of the present invention is operably linked to a regulatory sequence which is capable of providing for the expression of the coding sequence, such as  
15 by the chosen host cell. By way of example, the present invention covers a vector comprising the polynucleotide of the present invention operably linked to such a regulatory sequence, i.e. the vector is an expression vector.

The term "operably linked" refers to a juxtaposition wherein the components described are  
20 in a relationship permitting them to function in their intended manner. A regulatory sequence "operably linked" to a coding sequence is ligated in such a way that expression of the coding sequence is achieved under condition compatible with the control sequences.

The term "regulatory sequences" includes promoters and enhancers and other expression  
25 regulation signals.

The term "promoter" is used in the normal sense of the art, e.g. an RNA polymerase binding site.

Enhanced expression of the polynucleotide encoding the polypeptide of the present  
30 invention may also be achieved by the selection of heterologous regulatory regions, e.g. promoter, secretion leader and terminator regions, which serve to increase expression and, if desired, secretion levels of the protein of interest from the chosen expression host and/or

to provide for the inducible control of the expression of the polypeptide of the present invention

Preferably, the nucleotide sequence of the present invention may be operably linked to at least a promoter.

Aside from the promoter native to the gene encoding the polypeptide of the present invention, other promoters may be used to direct expression of the polypeptide of the present invention. The promoter may be selected for its efficiency in directing the expression of the polypeptide of the present invention in the desired expression host.

In another embodiment, a constitutive promoter may be selected to direct the expression of the desired polypeptide of the present invention. Such an expression construct may provide additional advantages since it circumvents the need to culture the expression hosts on a medium containing an inducing substrate.

Examples of strong constitutive and/or inducible promoters which are preferred for use in fungal expression hosts are those which are obtainable from the fungal genes for xylanase (*xlnA*), phytase, ATP-synthetase, subunit 9 (*oliC*), triose phosphate isomerase (*tpi*), alcohol dehydrogenase (*AdhA*),  $\alpha$ -amylase (*amy*), amyloglucosidase (AG - from the *glaA* gene), acetamidase (*amdS*) and glyceraldehyde-3-phosphate dehydrogenase (*gpd*) promoters.

Examples of strong yeast promoters are those obtainable from the genes for alcohol dehydrogenase, lactase, 3-phosphoglycerate kinase and triosephosphate isomerase.

Examples of strong bacterial promoters are the  $\alpha$ -amylase and *SP02* promoters as well as promoters from extracellular protease genes.

Hybrid promoters may also be used to improve inducible regulation of the expression construct.

The promoter can additionally include features to ensure or to increase expression in a suitable host. For example, the features can be conserved regions such as a Pribnow Box or a

TATA box. The promoter may even contain other sequences to affect (such as to maintain, enhance, decrease) the levels of expression of the nucleotide sequence of the present invention. For example, suitable other sequences include the Sh1-intron or an ADH intron. Other sequences include inducible elements - such as temperature, chemical, light or stress inducible elements. Also, suitable elements to enhance transcription or translation may be present. An example of the latter element is the TMV 5' signal sequence (see Sleat Gene 217 [1987] 217-225; and Dawson Plant Mol. Biol. 23 [1993] 97).

## SECRETION

Often, it is desirable for the polypeptide of the present invention to be secreted from the expression host into the culture medium from where the polypeptide of the present invention may be more easily recovered. According to the present invention, the secretion leader sequence may be selected on the basis of the desired expression host. Hybrid signal sequences may also be used with the context of the present invention.

Typical examples of heterologous secretion leader sequences are those originating from the fungal amyloglucosidase (AG) gene (*glaA* - both 18 and 24 amino acid versions e.g. from *Aspergillus*), the a-factor gene (yeasts e.g. *Saccharomyces* and *Kluyveromyces*) or the  $\alpha$ -amylase gene (*Bacillus*).

## CONSTRUCTS

The term "construct" - which is synonymous with terms such as "conjugate", "cassette" and "hybrid" - includes the nucleotide sequence according to the present invention directly or indirectly attached to a promoter. An example of an indirect attachment is the provision of a suitable spacer group such as an intron sequence, such as the Sh1-intron or the ADH intron, intermediate the promoter and the nucleotide sequence of the present invention. The same is true for the term "fused" in relation to the present invention which includes direct or indirect attachment. In each case, the terms do not cover the natural combination of the nucleotide sequence coding for the protein ordinarily associated with the wild type gene promoter and when they are both in their natural environment.



The construct may even contain or express a marker which allows for the selection of the genetic construct in, for example, a bacterium, preferably of the genus *Bacillus*, such as *Bacillus subtilis*, or plants into which it has been transferred. Various markers exist which may be used, such as for example those encoding mannose-6-phosphate isomerase (especially  
5 for plants) or those markers that provide for antibiotic resistance - e.g. resistance to G418, hygromycin, bleomycin, kanamycin and gentamycin.

Preferably the construct of the present invention comprises at least the nucleotide sequence of the present invention operably linked to a promoter.

10

## VECTORS

The term "vector" includes expression vectors and transformation vectors and shuttle vectors.

15 The term "expression vector" means a construct capable of *in vivo* or *in vitro* expression.

The term "transformation vector" means a construct capable of being transferred from one entity to another entity - which may be of the species or may be of a different species. If the construct is capable of being transferred from one species to another - such as from an *E.coli*  
20 plasmid to a bacterium, such as of the genus *Bacillus*, then the transformation vector is sometimes called a "shuttle vector". It may even be a construct capable of being transferred from an *E.coli* plasmid to an *Agrobacterium* to a plant.

The vectors of the present invention may be transformed into a suitable host cell as  
25 described below to provide for expression of a polypeptide of the present invention. Thus, in a further aspect the invention provides a process for preparing polypeptides according to the present invention which comprises cultivating a host cell transformed or transfected with an expression vector as described above under conditions to provide for expression by the vector of a coding sequence encoding the polypeptides, and recovering the expressed  
30 polypeptides.

The vectors may be for example, plasmid, virus or phage vectors provided with an origin of replication, optionally a promoter for the expression of the said polynucleotide and optionally a regulator of the promoter.

- 5 The vectors of the present invention may contain one or more selectable marker genes. The most suitable selection systems for industrial micro-organisms are those formed by the group of selection markers which do not require a mutation in the host organism. Examples of fungal selection markers are the genes for acetamidase (*amdS*), ATP synthetase, subunit 9 (*oliC*), orotidine-5'-phosphate-decarboxylase (*pvrA*), phleomycin and
- 10 benomyl resistance (*benA*). Examples of non-fungal selection markers are the bacterial G418 resistance gene (this may also be used in yeast, but not in filamentous fungi), the ampicillin resistance gene (*E. coli*), the neomycin resistance gene (*Bacillus*) and the *E.coli uidA* gene, coding for  $\beta$ -glucuronidase (GUS).

- 15 Vectors may be used *in vitro*, for example for the production of RNA or used to transfect or transform a host cell.

- Thus, polynucleotides of the present invention can be incorporated into a recombinant vector (typically a replicable vector), for example a cloning or expression vector. The
- 20 vector may be used to replicate the nucleic acid in a compatible host cell. Thus in a further embodiment, the invention provides a method of making polynucleotides of the present invention by introducing a polynucleotide of the present invention into a replicable vector, introducing the vector into a compatible host cell, and growing the host cell under conditions which bring about replication of the vector. The vector may be recovered from
- 25 the host cell. Suitable host cells are described below in connection with expression vectors.

- The present invention also relates to the use of genetically engineered host cells expressing a PDE\_XIV or variant, homologue, fragment or derivative thereof in screening methods for
- 30 the identification of inhibitors and antagonists of the PDE\_XIV that would modulate phosphodiesterase activity thereby modulating cyclic nucleotide levels. Such genetically engineered host cells could be used to screen peptide libraries or organic molecules capable of modulating PDE\_XIV activity. Antagonists and inhibitors of PDE\_XIV, such as

antibodies, peptides or small organic molecules will provide the basis for pharmaceutical compositions for the treatment of diseases associated with, for example, PDE\_XIV. Such inhibitors or antagonists can be administered alone or in combination with other therapeutics for the treatment of such diseases.

5

The present invention also relates to expression vectors and host cells comprising polynucleotide sequences encoding PDE\_XIV or variant, homologue, fragment or derivative thereof for the *in vivo* or *in vitro* production of PDE\_XIV protein or to screen for agents that can affect PDE\_XIV expression or activity.

10

#### TISSUE

The term "tissue" as used herein includes tissue *per se* and organ.

15 

#### HOST CELLS

The term "host cell" - in relation to the present invention includes any cell that could comprise the nucleotide sequence coding for the recombinant protein according to the present invention and/or products obtained therefrom, wherein a promoter can allow expression of the nucleotide sequence according to the present invention when present in the host cell.

20

Thus, a further embodiment of the present invention provides host cells transformed or transfected with a polynucleotide of the present invention. Preferably said polynucleotide is carried in a vector for the replication and expression of said polynucleotides. The cells will be chosen to be compatible with the said vector and may for example be prokaryotic (for example bacterial), fungal, yeast or plant cells.

25

The gram-negative bacterium *E. coli* is widely used as a host for heterologous gene expression. However, large amounts of heterologous protein tend to accumulate inside the cell. Subsequent purification of the desired protein from the bulk of *E.coli* intracellular proteins can sometimes be difficult.

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In contrast to *E.coli*, bacteria from the genus *Bacillus* are very suitable as heterologous hosts because of their capability to secrete proteins into the culture medium. Other bacteria suitable as hosts are those from the genera *Streptomyces* and *Pseudomonas*.

- 5 Depending on the nature of the polynucleotide encoding the polypeptide of the present invention, and/or the desirability for further processing of the expressed protein, eukaryotic hosts such as yeasts or other fungi may be preferred. In general, yeast cells are preferred over fungal cells because they are easier to manipulate. However, some proteins are either poorly secreted from the yeast cell, or in some cases are not processed properly (e.g. hyperglycosylation in yeast). In these instances, a different fungal host organism should be selected.

Examples of suitable expression hosts within the scope of the present invention are fungi such as *Aspergillus* species (such as those described in EP-A-0184438 and EP-A-0284603) and *Trichoderma* species; bacteria such as *Bacillus* species (such as those described in EP-A-0134048 and EP-A-0253455), *Streptomyces* species and *Pseudomonas* species; and yeasts such as *Kluyveromyces* species (such as those described in EP-A-0096430 and EP-A-0301670) and *Saccharomyces* species. By way of example, typical expression hosts may be selected from *Aspergillus niger*, *Aspergillus niger* var. *tubigenis*, *Aspergillus niger* var. *awamori*, *Aspergillus aculeatis*, *Aspergillus nidulans*, *Aspergillus oryzae*, *Trichoderma reesei*, *Bacillus subtilis*, *Bacillus licheniformis*, *Bacillus amyloliquefaciens*, *Kluyveromyces lactis* and *Saccharomyces cerevisiae*.

The use of suitable host cells - such as yeast, fungal and plant host cells - may provide for post-translational modifications (e.g. myristoylation, glycosylation, truncation, lipidation and tyrosine, serine or threonine phosphorylation) as may be needed to confer optimal biological activity on recombinant expression products of the present invention.

## ORGANISM

30

The term "organism" in relation to the present invention includes any organism that could comprise the nucleotide sequence coding for the recombinant protein according to the present invention and/or products obtained therefrom, wherein a promoter can allow expression of

the nucleotide sequence according to the present invention when present in the organism.

Examples of organisms may include a fungus, yeast or a plant.

The term "transgenic organism" in relation to the present invention includes any organism  
5 that comprises the nucleotide sequence coding for the protein according to the present  
invention and/or products obtained therefrom, wherein the promoter can allow expression of  
the nucleotide sequence according to the present invention within the organism. Preferably  
the nucleotide sequence is incorporated in the genome of the organism.

10 The term "transgenic organism" does not cover the native nucleotide coding sequence  
according to the present invention in its natural environment when it is under the control of  
its native promoter which is also in its natural environment. In addition, the present invention  
does not cover the native protein according to the present invention when it is in its natural  
environment and when it has been expressed by its native nucleotide coding sequence which  
15 is also in its natural environment and when that nucleotide sequence is under the control of its  
native promoter which is also in its natural environment.

Therefore, the transgenic organism of the present invention includes an organism comprising  
any one of, or combinations of, the nucleotide sequence coding for the amino acid sequence  
20 according to the present invention, constructs according to the present invention (including  
combinations thereof), vectors according to the present invention, plasmids according to the  
present invention, cells according to the present invention, tissues according to the present  
invention or the products thereof. The transformed cell or organism could prepare acceptable  
quantities of the desired compound which would be easily retrievable from, the cell or  
25 organism.

The present invention also encompasses organisms that have been modified so that they  
express elevated levels of the enzyme.

## ATTENUATION/KNOCK-OUT ANIMAL MODELS

The present invention also encompasses organisms that have been modified so that the enzyme activity has been attenuated or even eliminated. An example of such an embodiment is a knock-out rat or mouse wherein the natural coding sequence has been removed and replaced with, for example, a reporter gene – such as  $\beta$ -gal. Alternatively, the promoter can be silenced so that no gene expression occurs. These types of organisms may be prepared by use of standard techniques.

## 10 TRANSFORMATION OF HOST CELLS/HOST ORGANISMS

As indicated earlier, the host organism can be a prokaryotic or a eukaryotic organism. Examples of suitable prokaryotic hosts include *E. coli* and *Bacillus subtilis*. Teachings on the transformation of prokaryotic hosts is well documented in the art, for example see Sambrook et al (Molecular Cloning: A Laboratory Manual, 2nd edition, 1989, Cold Spring Harbor Laboratory Press) and Ausubel *et al.*, Current Protocols in Molecular Biology (1995), John Wiley & Sons, Inc.

If a prokaryotic host is used then the nucleotide sequence may need to be suitably modified before transformation - such as by removal of introns.

In another embodiment the transgenic organism can be a yeast. In this regard, yeast have also been widely used as a vehicle for heterologous gene expression. The species *Saccharomyces cerevisiae* has a long history of industrial use, including its use for heterologous gene expression. Expression of heterologous genes in *Saccharomyces cerevisiae* has been reviewed by Goodey et al (1987, Yeast Biotechnology, D R Berry et al, eds, pp 401-429, Allen and Unwin, London) and by King et al (1989, Molecular and Cell Biology of Yeasts, E F Walton and G T Yarronton, eds, pp 107-133, Blackie, Glasgow).

For several reasons *Saccharomyces cerevisiae* is well suited for heterologous gene expression. First, it is non-pathogenic to humans and it is incapable of producing certain endotoxins. Second, it has a long history of safe use following centuries of commercial exploitation for various purposes. This has led to wide public acceptability. Third, the

extensive commercial use and research devoted to the organism has resulted in a wealth of knowledge about the genetics and physiology as well as large-scale fermentation characteristics of *Saccharomyces cerevisiae*.

- 5 A review of the principles of heterologous gene expression in *Saccharomyces cerevisiae* and secretion of gene products is given by E Hinchcliffe E Kenny (1993, "Yeast as a vehicle for the expression of heterologous genes", Yeasts, Vol 5, Anthony H Rose and J Stuart Harrison, eds, 2nd edition, Academic Press Ltd.).

- 10 Several types of yeast vectors are available, including integrative vectors, which require recombination with the host genome for their maintenance, and autonomously replicating plasmid vectors.

- In order to prepare the transgenic *Saccharomyces*, expression constructs are prepared by  
15 inserting the nucleotide sequence of the present invention into a construct designed for expression in yeast. Several types of constructs used for heterologous expression have been developed. The constructs contain a promoter active in yeast fused to the nucleotide sequence of the present invention, usually a promoter of yeast origin, such as the GAL1 promoter, is used. Usually a signal sequence of yeast origin, such as the sequence encoding  
20 the SUC2 signal peptide, is used. A terminator active in yeast ends the expression system.

- For the transformation of yeast several transformation protocols have been developed. For example, a transgenic *Saccharomyces* according to the present invention can be prepared by following the teachings of Hinnen et al (1978, Proceedings of the National Academy of  
25 Sciences of the USA 75, 1929); Beggs, J D (1978, Nature, London, 275, 104); and Ito, H et al (1983, J Bacteriology 153, 163-168).

- The transformed yeast cells are selected using various selective markers. Among the markers used for transformation are a number of auxotrophic markers such as LEU2, HIS4 and TRP1,  
30 and dominant antibiotic resistance markers such as aminoglycoside antibiotic markers, eg G418.

Another host organism is a plant. The basic principle in the construction of genetically modified plants is to insert genetic information in the plant genome so as to obtain a stable maintenance of the inserted genetic material.

- 5 Several techniques exist for inserting the genetic information, the two main principles being direct introduction of the genetic information and introduction of the genetic information by use of a vector system. A review of the general techniques may be found in articles by Potrykus (*Annu Rev Plant Physiol Plant Mol Biol* [1991] 42:205-225) and Christou (*Agro-Food-Industry Hi-Tech* March/April 1994 17-27). Further teachings on plant transformation  
10 may be found in EP-A-0449375.

Thus, the present invention also provides a method of transforming a host cell with a nucleotide sequence shown as any one of the sequences shown in the attached sequence listings or a derivative, homologue, variant or fragment thereof.

- 15 Host cells transformed with a PDE nucleotide coding sequence may be cultured under conditions suitable for the expression and recovery of the encoded protein from cell culture. The protein produced by a recombinant cell may be secreted or may be contained intracellularly depending on the sequence and/or the vector used. As will be understood by  
20 those of skill in the art, expression vectors containing PDE coding sequences can be designed with signal sequences which direct secretion of PDE coding sequences through a particular prokaryotic or eukaryotic cell membrane. Other recombinant constructions may join PDE coding sequence to nucleotide sequence encoding a polypeptide domain which will facilitate purification of soluble proteins (Kroll *DJ et al* (1993) *DNA Cell Biol* 12:441-  
25 53, see also above discussion of vectors containing fusion proteins).

## PRODUCTION OF THE POLYPEPTIDE

- According to the present invention, the production of the polypeptide of the present  
30 invention can be effected by the culturing of, for example, microbial expression hosts, which have been transformed with one or more polynucleotides of the present invention, in a conventional nutrient fermentation medium. The selection of the appropriate medium may be based on the choice of expression hosts and/or based on the regulatory



requirements of the expression construct. Such media are well-known to those skilled in the art. The medium may, if desired, contain additional components favouring the transformed expression hosts over other potentially contaminating micro-organisms.

- 5 Thus, the present invention also provides a method for producing a polypeptide having PDE\_XIV activity, the method comprising the steps of a) transforming a host cell with a nucleotide sequence shown as any one of the sequences shown in the attached sequence listings or a derivative, homologue, variant or fragment thereof; and b) culturing the transformed host cell under conditions suitable for the expression of said polypeptide.

10

The present invention also provides a method for producing a polypeptide having PDE\_XIV activity, the method comprising the steps of a) culturing a host cell that has been transformed with a nucleotide sequence shown as any one of the sequences shown in the attached sequence listings or a derivative, homologue, variant or fragment thereof under  
15 conditions suitable for the expression of said polypeptide; and b) recovering said polypeptide from the host cell culture.

20

The present invention also provides a method for producing a polypeptide having PDE\_XIV activity, the method comprising the steps of a) transforming a host cell with a nucleotide sequence shown as any one of the sequences shown in the attached sequence listings or a derivative, homologue, variant or fragment thereof; b) culturing the transformed host cell under conditions suitable for the expression of said polypeptide; and  
c) recovering said polypeptide from the host cell culture.

25

## RIBOZYMES

Ribozymes are enzymatic RNA molecules capable of catalysing the specific cleavage of RNA. The mechanism of ribozyme action involves sequence specific hybridisation of the ribozyme molecule to complementary target RNA, followed by an endonucleolytic cleavage.

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Within the scope of the invention are engineered hammerhead motif ribozyme molecules that specifically and efficiently catalyse endonucleolytic cleavage of PDE RNA sequences.

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Specific ribozyme cleavage sites within any potential RNA target are initially identified by scanning the target molecule for ribozyme cleavage sites which include the following sequences, GUA, GUU and GUC. Once identified, short RNA sequences of between 15 and 20 ribonucleotides corresponding to the region of the target gene containing the cleavage site may be evaluated for secondary structural features which may render the oligonucleotide sequence inoperable. The suitability of candidate targets may also be evaluated by testing accessibility to hybridisation with complementary oligonucleotides using ribonuclease protection assays.

- Both antisense RNA and DNA molecules and ribozymes of the invention may be prepared by any method known in the art for the synthesis of RNA molecules. These include techniques for chemically synthesising oligonucleotides such as solid phase phosphoramidite chemical synthesis. Alternatively, RNA molecules may be generated by *in vitro* or *in vivo* transcription of DNA sequences encoding the antisense RNA molecule.
- Such DNA sequences may be incorporated into a wide variety of vectors with suitable RNA polymerase promoters such as T7 or SP6. Alternatively, antisense cDNA constructs that synthesize antisense RNA constitutively or inducibly can be introduced into cell lines, cells or tissues.

## DETECTION

- The presence of the PDE polynucleotide coding sequence can be detected by DNA-DNA or DNA-RNA hybridisation or amplification using probes, portions or fragments of the sequence presented as any one of the sequences shown in the attached sequence listings.
- Nucleic acid amplification based assays involve the use of oligonucleotides or oligomers based on the PDE coding sequence to detect transformants containing PDE DNA or RNA. As used herein "oligonucleotides" or "oligomers" may refer to a nucleic acid sequence of at least about 10 nucleotides and as many as about 60 nucleotides, preferably about 15 to 30 nucleotides, and more preferably about 20-25 nucleotides which can be used as a probe or primer. Preferably, oligonucleotides are derived from the 3' region of the nucleotide sequence shown as any one of the sequences shown in the attached sequence listings.

A variety of protocols for detecting and measuring the expression of PDE polypeptide, such as by using either polyclonal or monoclonal antibodies specific for the protein, are known in the art. Examples include enzyme-linked immunosorbent assay (ELISA), radioimmunoassay (RIA) and fluorescent activated cell sorting (FACS). A two-site, monoclonal-based immunoassay utilising monoclonal antibodies reactive to two non-interfering epitopes on PDE polypeptides is preferred, but a competitive binding assay may be employed. These and other assays are described, among other places, in Hampton R *et al* (1990, Serological Methods, A Laboratory Manual, APS Press, St Paul MN) and Maddox DE *et al* (1983, J Exp Med 15 8:121 1).

A wide variety of labels and conjugation techniques are known by those skilled in the art and can be used in various nucleic and amino acid assays. Means for producing labelled hybridisation or PCR probes for detecting PDE polynucleotide sequences include oligolabelling, nick translation, end-labelling or PCR amplification using a labelled nucleotide. Alternatively, the PDE coding sequence, or any portion of it, may be cloned into a vector for the production of an mRNA probe. Such vectors are known in the art, are commercially available, and may be used to synthesize RNA probes *in vitro* by addition of an appropriate RNA polymerase such as T7, T3 or SP6 and labelled nucleotides.

A number of companies such as Pharmacia Biotech (Piscataway, NJ), Promega (Madison, WI), and US Biochemical Corp (Cleveland, OH) supply commercial kits and protocols for these procedures. Suitable reporter molecules or labels include those radionuclides, enzymes, fluorescent, chemiluminescent, or chromogenic agents as well as substrates, cofactors, inhibitors, magnetic particles and the like. Patents teaching the use of such labels include US-A-3817837; US-A-3850752; US-A-3939350; US-A-3996345; US-A-4277437; US-A-4275149 and US-A-4366241. Also, recombinant immunoglobulins may be produced as shown in US-A-4816567.

Additional methods to quantify the expression of a particular molecule include radiolabeling (Melby PC *et al* 1993 J Immunol Methods 159:235-44) or biotinylating (Duplaa C *et al* 1993 Anal Biochem 229:36) nucleotides, coamplification of a control nucleic acid, and standard curves onto which the experimental results are interpolated. Quantification of multiple samples may be speeded up by running the assay in an ELISA

format where the oligomer of interest is presented in various dilutions and a spectrophotometric or calorimetric response gives rapid quantification.

Although the presence/absence of marker gene expression suggests that the gene of interest is also present, its presence and expression should be confirmed. For example, if the PDE coding sequence is inserted within a marker gene sequence, recombinant cells containing PDE coding regions can be identified by the absence of marker gene function. Alternatively, a marker gene can be placed in tandem with a PDE coding sequence under the control of a single promoter. Expression of the marker gene in response to induction or selection usually indicates expression of PDE as well.

Alternatively, host cells which contain the coding sequence for PDE and express PDE coding regions may be identified by a variety of procedures known to those of skill in the art. These procedures include, but are not limited to, DNA-DNA or DNA-RNA hybridisation and protein bioassay or immunoassay techniques which include membrane-based, solution-based, or chip-based technologies for the detection and/or quantification of the nucleic acid or protein.

## ANTIBODIES

The amino acid sequence of the present invention can also be used to generate antibodies - such as by use of standard techniques - against the amino acid sequence.

Procedures well known in the art may be used for the production of antibodies to PDE\_XIV polypeptides. Such antibodies include, but are not limited to, polyclonal, monoclonal, chimeric, single chain, Fab fragments and fragments produced by a Fab expression library. Neutralising antibodies, i.e., those which inhibit biological activity of PDE polypeptides, are especially preferred for diagnostics and therapeutics.

For the production of antibodies, various hosts including goats, rabbits, rats, mice, etc. may be immunised by injection with the inhibitor or any portion, variant, homologue, fragment or derivative thereof or oligopeptide which retains immunogenic properties. Depending on the host species, various adjuvants may be used to increase immunological response. Such

adjuvants include, but are not limited to, Freund's, mineral gels such as aluminium hydroxide, and surface active substances such as lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet hemocyanin, and dinitrophenol. BCG (*Bacilli Calmette-Guerin*) and *Corynebacterium parvum* are potentially useful human adjuvants which may be employed.

Monoclonal antibodies to the amino acid sequence may be even prepared using any technique which provides for the production of antibody molecules by continuous cell lines in culture. These include, but are not limited to, the hybridoma technique originally described by Koehler and Milstein (1975 Nature 256:495-497), the human B-cell hybridoma technique (Kosbor *et al* (1983) Immunol Today 4:72; Cote *et al* (1983) Proc Natl Acad Sci 80:2026-2030) and the EBV-hybridoma technique (Cole *et al* (1985) Monoclonal Antibodies and Cancer Therapy, Alan R Liss Inc, pp 77-96). In addition, techniques developed for the production of "chimeric antibodies", the splicing of mouse antibody genes to human antibody genes to obtain a molecule with appropriate antigen specificity and biological activity can be used (Morrison *et al* (1984) Proc Natl Acad Sci 81:6851-6855; Neuberger *et al* (1984) Nature 312:604-608; Takeda *et al* (1985) Nature 314:452-454). Alternatively, techniques described for the production of single chain antibodies (US-A-4946779) can be adapted to produce inhibitor specific single chain antibodies.

Antibodies may also be produced by inducing *in vivo* production in the lymphocyte population or by screening recombinant immunoglobulin libraries or panels of highly specific binding reagents as disclosed in Orlandi *et al* (1989, Proc Natl Acad Sci 86: 3833-3837), and Winter G and Milstein C (1991; Nature 349:293-299).

Antibody fragments which contain specific binding sites for PDE<sub>XIV</sub> may also be generated. For example, such fragments include, but are not limited to, the F(ab')<sub>2</sub> fragments which can be produced by pepsin digestion of the antibody molecule and the Fab fragments which can be generated by reducing the disulphide bridges of the F(ab')<sub>2</sub> fragments. Alternatively, Fab expression libraries may be constructed to allow rapid and easy identification of monoclonal Fab fragments with the desired specificity (Huse WD *et al* (1989) Science 256:1275-1281).

An alternative technique involves screening phage display libraries where, for example the phage express scFv fragments on the surface of their coat with a large variety of complementarity determining regions (CDRs). This technique is well known in the art.

5

PDE\_XIV-specific antibodies are useful for the diagnosis of conditions and diseases associated with expression of PDE\_XIV polypeptide. A variety of protocols for competitive binding or immunoradiometric assays using either polyclonal or monoclonal antibodies with established specificities are well known in the art. Such immunoassays typically involve the formation of complexes between PDE\_XIV polypeptides and its specific antibody (or similar PDE\_XIV-binding molecule) and the measurement of complex formation. A two-site, monoclonal based immunoassay utilising monoclonal antibodies reactive to two non-interfering epitopes on a specific PDE\_XIV protein is preferred, but a competitive binding assay may also be employed. These assays are described in Maddox DE *et al* (1983, J Exp Med 158:121 1).

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15

Anti-PDE\_XIV antibodies are useful for the diagnosis of inflammation, conditions associated with proliferation of hematopoietic cells and HIV infection or other disorders or diseases characterised by abnormal expression of a PDE\_XIV. Diagnostic assays for a PDE\_XIV include methods utilising the antibody and a label to detect a PDE\_XIV polypeptide in human body fluids, cells, tissues or sections or extracts of such tissues. The polypeptides and antibodies of the present invention may be used with or without modification. Frequently, the polypeptides and antibodies will be labelled by joining them, either covalently or noncovalently, with a reporter molecule. A wide variety of reporter molecules are known to those of skill in the art.

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Antibodies may be used in method of detecting polypeptides of the invention present in biological samples by a method which comprises: (a) providing an antibody of the invention; (b) incubating a biological sample with said antibody under conditions which allow for the formation of an antibody-antigen complex; and (c) determining whether antibody-antigen complex comprising said antibody is formed.

30

Depending on the circumstances, suitable samples may include extracts tissues such as brain, breast, ovary, lung, colon, pancreas, testes, liver, muscle and bone tissues or from neoplastic growths derived from such tissues.

- 5     Antibodies of the invention may be bound to a solid support and/or packaged into kits in a suitable container along with suitable reagents, controls, instructions and the like.

#### ASSAYS/IDENTIFICATION METHODS

- 10     The present invention also relates to an assay method for detecting the presence of PDE\_XIV in cells (such as human cells) comprising: (a) performing a reverse transcriptase-polymerase chain reaction on RNA (such as total RNA) from such cells using a pair of polymerase chain reaction primers that are specific for PDE\_XIV, as determined from the DNA sequence of  
15     thereof; and (b) assaying the appearance of an appropriately sized PCR (polymerase chain reaction) fragment - such as by agarose gel electrophoresis.

- The present invention also relates to a method of identifying agents (such as compounds, other substances or compositions comprising same) that affect (such as inhibit or otherwise  
20     modify) the activity of PDE\_XIV and/or the expression thereof, the method comprising contacting PDE\_XIV or the nucleotide sequence coding for same with the agent and then measuring the activity of PDE\_XIV and/or the expression thereof.

- The present invention also relates to a method of identifying agents (such as compounds,  
25     other substances or compositions comprising same) that selectively affect (such as inhibit or otherwise modify) the activity of PDE\_XIV and/or the expression thereof, the method comprising contacting PDE\_XIV or the nucleotide sequence coding for same with the agent and then measuring the activity of PDE\_XIV and/or the expression thereof.

- 30     The present invention also relates to a method of identifying agents (such as compounds, other substances or compositions comprising same) that selectively affect (such as inhibit or otherwise modify) the activity of PDE\_XIVA1 or PDE\_XIVA2 or PDE\_XIVA3 or PDE\_XIVA4 and/or the expression thereof, the method comprising contacting the relevant

PDE\_XIV or the nucleotide sequence coding for same with the agent and then measuring the activity of the relevant PDE\_XIV and/or the expression thereof.

5 The present invention also relates to a method of identifying agents (such as compounds, other substances or compositions comprising same) that affect (such as inhibit or otherwise modify) the activity of PDE\_XIV and/or the expression thereof, the method comprising measuring the activity of PDE\_XIV and/or the expression thereof in the presence of the agent or after the addition of the agent in: (a) a cell line into which has been incorporated recombinant DNA comprising the DNA sequence of any one of the sequences shown in the  
10 attached sequence listings or an allelic variation thereof, or (b) a cell population or cell line that naturally selectively expresses PDE\_XIV. Preferably, the activity of PDE\_XIV is determined by the assay method described above.

15 The present invention also relates to a method of identifying agents (such as compounds, other substances or compositions comprising same) that selectively affect (such as inhibit or otherwise modify) the activity of PDE\_XIV and/or the expression thereof, the method comprising measuring the activity of PDE\_XIV and/or the expression thereof in the presence of the agent or after the addition of the agent in: (a) a cell line into which has been incorporated recombinant DNA comprising the DNA sequence of any one of the sequences  
20 shown in the attached sequence listings or an allelic variation thereof, or (b) a cell population or cell line that naturally selectively expresses PDE\_XIV. Preferably, the activity of PDE\_XIV is determined by the assay method described above.

25 The present invention also relates to a method of identifying agents (such as compounds, other substances or compositions comprising same) that selectively affect (such as inhibit or otherwise modify) the activity of PDE\_XIVA1 or PDE\_XIVA2 or PDE\_XIVA3 or PDE\_XIVA4 and/or the expression thereof, the method comprising measuring the activity of the respective PDE\_XIV and/or the expression thereof in the presence of the agent or after the addition of the agent in: (a) a cell line into which has been incorporated recombinant  
30 DNA comprising the respective DNA sequence of any one of the sequences shown in the attached sequence listings or an allelic variation thereof, or (b) a cell population or cell line that naturally selectively expresses the respective PDE\_XIV. Preferably, the activity of PDE\_XIV is determined by the assay method described above.



The present invention also provides a method of screening an agent for modulation (preferably for specific modulation) of PDE\_XIV (or a derivative, homologue, variant or fragment thereof) activity or the expression of the nucleotide sequence coding for same (including a derivative, homologue, variant or fragment thereof), the method comprising the steps of: a) providing a candidate agent; b) combining PDE\_XIV (or the derivative, homologue, variant or fragment thereof) or the nucleotide sequence coding for same (or the derivative, homologue, variant or fragment thereof) with the candidate agent for a time sufficient to allow modulation under suitable conditions; and c) detecting modulation of the candidate agent to PDE\_XIV (or the derivative, homologue, variant or fragment thereof) or the nucleotide sequence coding for same (or the derivative, homologue, variant or fragment thereof) in order to ascertain if the candidate agent modulates PDE\_XIV (or the derivative, homologue, variant or fragment thereof) activity or the expression of the nucleotide sequence coding for same (or the derivative, homologue, variant or fragment thereof).

The present invention also provides a method of screening an agent for specific binding affinity with PDE\_XIV (or a derivative, homologue, variant or fragment thereof) or the nucleotide sequence coding for same (including a derivative, homologue, variant or fragment thereof), the method comprising the steps of: a) providing a candidate agent; b) combining PDE\_XIV (or the derivative, homologue, variant or fragment thereof) or the nucleotide sequence coding for same (or the derivative, homologue, variant or fragment thereof) with the candidate agent for a time sufficient to allow binding under suitable conditions; and c) detecting binding of the candidate agent to PDE\_XIV (or the derivative, homologue, variant or fragment thereof) or the nucleotide sequence coding for same (or the derivative, homologue, variant or fragment thereof) in order to ascertain if the candidate agent binds to PDE\_XIV (or the derivative, homologue, variant or fragment thereof) or the nucleotide sequence coding for same (or the derivative, homologue, variant or fragment thereof).

30

The present invention also provides a method of identifying an agent which is capable of modulating PDE\_XIV, the method comprising the steps of: a) contacting the agent with PDE\_XIV (or a derivative, homologue, variant or fragment thereof) or the nucleotide

sequence coding for same (or the derivative, homologue, variant or fragment thereof), b) incubating the mixture of step a) with a cyclic nucleotide under conditions suitable for the hydrolysis of the cyclic nucleotide, c) measuring the amount of cyclic nucleotide hydrolysis, and d) comparing the amount of cyclic nucleotide hydrolysis of step c) with the amount of cyclic nucleotide hydrolysis obtained with PDE\_XIV (or the derivative, homologue, variant or fragment thereof) or the nucleotide sequence coding for same (or the derivative, homologue, variant or fragment thereof) incubated without the compound, thereby determining whether the agent affects (such as stimulates or inhibits) cyclic nucleotide hydrolysis.

10

Thus, in certain embodiments of the present invention, PDE\_XIV or a variant, homologue, fragment or derivative thereof and/or a cell line that expresses the PDE\_XIV or variant, homologue, fragment or derivative thereof may be used to screen for antibodies, peptides, or other agent, such as organic or inorganic molecules, that act as modulators of phosphodiesterase activity or for the expression thereof, thereby identifying a therapeutic agent capable of modulating cyclic nucleotide levels. For example, anti-PDE\_XIV antibodies capable of neutralising the activity of PDE\_XIV may be used to inhibit PDE\_XIV hydrolysis of cyclic nucleotides, thereby increasing their levels. Alternatively, screening of peptide libraries or organic libraries made by combinatorial chemistry with recombinantly expressed PDE\_XIV or a variant, homologue, fragment or derivative thereof or cell lines expressing PDE\_XIV or a variant, homologue, fragment or derivative thereof may be useful for identification of therapeutic agents that function by modulating PDE\_XIV hydrolysis of cyclic nucleotides. Synthetic compounds, natural products, and other sources of potentially biologically active materials can be screened in a number of ways deemed to be routine to those of skill in the art. For example, nucleotide sequences encoding the N-terminal region of PDE\_XIV may be expressed in a cell line which can be used for screening of allosteric modulators, either agonists or antagonists, of PDE\_XIV activity. Alternatively, nucleotide sequences encoding the conserved catalytic domain of PDE\_XIV can be expressed in cell lines and used to screen for inhibitors of cyclic nucleotide hydrolysis.

30

The ability of a test agent to interfere with PDE\_XIV activity or cyclic nucleotide hydrolysis may be determined by measuring PDE\_XIV levels or cyclic nucleotide levels

(as disclosed in Smith *et al* 1993 Appl. Biochem. Biotechnol. 41:189-218). There are also commercially available immunoassay kits for the measurement of cAMP and cGMP (eg Amersham International, Arlington Heights, IL and DuPont, Boston, MA). The activity of PDE\_XIV may also be monitored by measuring other responses such as phosphorylation or dephosphorylation of other proteins using conventional techniques developed for these purpose.

Accordingly, the present invention provides a method of identifying a compound which is capable of modulating the cyclic nucleotide phosphodiesterase activity of a PDE\_XIV, or a variant, homologue, fragment or derivative thereof, comprising the steps of a) contacting the compound with a PDE\_XIV, or a variant, homologue, fragment or derivative thereof; b) incubating the mixture of step a) with a cyclic nucleotide under conditions suitable for the hydrolysis of the cyclic nucleotide; c) measuring the amount of cyclic nucleotide hydrolysis; and d) comparing the amount of cyclic nucleotide hydrolysis of step c) with the amount of cyclic nucleotide hydrolysis obtained with the PDE\_XIV, or a variant, homologue, fragment or derivative thereof, incubated without the compound, thereby determining whether the compound stimulates or inhibits cyclic nucleotide hydrolysis. In one embodiment of the method, the fragment may be from the N-terminal region of the PDE\_XIV and provides a method to identify allosteric modulators of the PDE\_XIV. In another embodiment of the present invention, the fragment may be from the carboxy terminal region of the PDE\_XIV and provides a method to identify inhibitors of cyclic nucleotide hydrolysis.

A PDE\_XIV polypeptide, its immunogenic fragments or oligopeptides thereof can be used for screening therapeutic compounds in any of a variety of drug screening techniques. The polypeptide employed in such a test may be free in solution, affixed to a solid support, borne on a cell surface, or located intracellularly. The abolition of activity or the formation of binding complexes between a PDE\_XIV polypeptide and the agent being tested may be measured.

Accordingly, the present invention provides a method for screening one or a plurality of compounds for modulation (preferably specific modulation, such as specific binding affinity) of PDE\_XIV or the expression thereof, or a portion thereof or variant, homologue,

fragment or derivative thereof, comprising providing one or a plurality of compounds; combining a PDE\_XIV or a nucleotide sequence coding for same or a portion thereof or variant, homologue, fragment or derivative thereof with the or each of a plurality of compounds for a time sufficient to allow modulation under suitable conditions; and  
5 detecting binding of a PDE\_XIV, or portion thereof or variant, homologue, fragment or derivative thereof, to each of the plurality of compounds, thereby identifying the compound or compounds which modulate a PDE\_XIV or a nucleotide sequence coding for same. In such an assay, the plurality of compounds may be produced by combinatorial chemistry techniques known to those of skill in the art.

10

Another technique for drug screening provides for high throughput screening of compounds having suitable binding affinity to the PDE\_XIV polypeptides and is based upon the method described in detail in Geysen, European Patent Application 84/03564, published on September 13, 1984. In summary, large numbers of different small peptide  
15 test compounds are synthesized on a solid substrate, such as plastic pins or some other surface. The peptide test compounds are reacted with PDE\_XIV fragments and washed. A bound PDE\_XIV is then detected - such as by appropriately adapting methods well known in the art. A purified PDE\_XIV can also be coated directly onto plates for use in the aforementioned drug screening techniques. Alternatively, non-neutralising antibodies can  
20 be used to capture the peptide and immobilise it on a solid support.

This invention also contemplates the use of competitive drug screening assays in which neutralising antibodies capable of binding a PDE\_XIV specifically compete with a test compound for binding a PDE\_XIV. In this manner, the antibodies can be used to detect the  
25 presence of any peptide which shares one or more antigenic determinants with a PDE\_XIV.

The assay method of the present invention may be a high throughput screen (HTS). In this regard, the teachings of WO 84/03564 may be adapted for the PDE of the present invention.

30

The teachings of US-A-5738985 may be adapted for the assay method of the present invention.

The present invention also provides one or more agents identified by the assays methods and identification methods of the present invention.

5

The agent of the present invention can be, for example, an organic compound or an inorganic compound. The agent can be, for example, a nucleotide sequence that is anti-sense to all or part of the sequences shown in the attached sequence listings.

10 The invention further provides an agent of the present invention (or even a pharmaceutically acceptable salt thereof, or a pharmaceutically acceptable solvate thereof) or a pharmaceutical composition containing any of the foregoing, for use as a medicament.

15 The present invention also provides the use of an agent to affect PDE\_XIV activity (such as to inhibit, modulate or agonise) in any one or more of the putamen, the Caudate nucleus of the brain, the Occipital lobe of the brain, the heart, ovary, the pituitary gland, kidney, liver, small intestine, thymus, skeletal muscle, leukocyte regions, dorsal root ganglia, uterus, cochlea, small intestine (duodenum), astrocytoma, and appendix.

20 **DIAGNOSTICS**

The present invention also provides a diagnostic composition for the detection of PDE\_XIV polynucleotide sequences. The diagnostic composition may comprise any one of the sequences shown in the attached sequence listings or a variant, homologue, fragment  
25 or derivative thereof, or a sequence capable of hybridising to all or part of any one of the sequences shown in the attached sequence listings or an allelic variation thereof.

In order to provide a basis for the diagnosis of disease, normal or standard values from a PDE\_XIV polypeptide expression should be established. This is accomplished by  
30 combining body fluids or cell extracts taken from normal subjects, either animal or human, with antibody to a PDE\_XIV polypeptide under conditions suitable for complex formation which are well known in the art. The amount of standard complex formation may be quantified by comparing it to a dilution series of positive controls where a known amount

of antibody is combined with known concentrations of a purified PDE\_XIV polypeptide. Then, standard values obtained from normal samples may be compared with values obtained from samples from subjects potentially affected by a disorder or disease related to a PDE\_XIV polypeptide expression. Deviation between standard and subject values establishes the presence of the disease state.

A PDE\_XIV polynucleotide, or any part thereof, may provide the basis for a diagnostic and/or a therapeutic compound. For diagnostic purposes, PDE\_XIV polynucleotide sequences may be used to detect and quantify gene expression in conditions, disorders or diseases in which PDE\_XIV activity may be implicated.

PDE\_XIV encoding polynucleotide sequence may be used for the diagnosis of diseases resulting from expression of PDE\_XIV. For example, polynucleotide sequences encoding PDE\_XIV may be used in hybridisation or PCR assays of tissues from biopsies or autopsies or biological fluids, such as serum, synovial fluid or tumour biopsy, to detect abnormalities in PDE\_XIV expression. The form of such qualitative or quantitative methods may include Southern or northern analysis, dot blot or other membrane-based technologies; PCR technologies; dip stick, pin or chip technologies; and ELISA or other multiple sample format technologies. All of these techniques are well known in the art and are in fact the basis of many commercially available diagnostic kits.

Such assays may be tailored to evaluate the efficacy of a particular therapeutic treatment regime and may be used in animal studies, in clinical trials, or in monitoring the treatment of an individual patient. In order to provide a basis for the diagnosis of disease, a normal or standard profile for PDE expression should be established. This is accomplished by combining body fluids or cell extracts taken from normal subjects, either animal or human, with PDE\_XIV or a portion thereof, under conditions suitable for hybridisation or amplification. Standard hybridisation may be quantified by comparing the values obtained for normal subjects with a dilution series of positive controls run in the same experiment where a known amount of purified PDE\_XIV is used. Standard values obtained from normal samples may be compared with values obtained from samples from subjects potentially affected by a disorder or disease related to expression of the PDE coding sequence. Deviation between standard and subject values establishes the presence of the

disease state. If disease is established, an existing therapeutic agent is administered, and treatment profile or values may be generated. Finally, the assay may be repeated on a regular basis to evaluate whether the values progress toward or return to the normal or standard pattern. Successive treatment profiles may be used to show the efficacy of treatment over a period of several days or several months.

Thus, the present invention relates to the use of a PDE\_XIV polypeptide, or variant, homologue, fragment or derivative thereof, to produce anti-PDE\_XIV antibodies which can, for example, be used diagnostically to detect and quantify PDE\_XIV levels in disease states.

The present invention further provides diagnostic assays and kits for the detection of PDE\_XIV in cells and tissues comprising a purified PDE\_XIV which may be used as a positive control, and anti-PDE\_XIV antibodies. Such antibodies may be used in solution-based, membrane-based, or tissue-based technologies to detect any disease state or condition related to the expression of PDE\_XIV protein or expression of deletions or a variant, homologue, fragment or derivative thereof.

## PROBES

Another aspect of the subject invention is the provision of nucleic acid hybridisation or PCR probes which are capable of detecting polynucleotide sequences, including genomic sequences, encoding PDE coding region or closely related molecules, such as alleles. The specificity of the probe, i.e., whether it is derived from a highly conserved, conserved or non-conserved region or domain, and the stringency of the hybridisation or amplification (high, intermediate or low) will determine whether the probe identifies only naturally occurring PDE coding sequence, or related sequences. Probes for the detection of related nucleic acid sequences are selected from conserved or highly conserved nucleotide regions of cyclic nucleotide PDE family members, such as the 3' region, and such probes may be used in a pool of degenerate probes. For the detection of identical nucleic acid sequences, or where maximum specificity is desired, nucleic acid probes are selected from the non-conserved nucleotide regions or unique regions of PDE polynucleotides. As used herein, the term "non-conserved nucleotide region" refers to a nucleotide region that is unique to

the PDE coding sequence disclosed herein and does not occur in related family members, such as known cyclic nucleotide PDEs.

PCR as described in US-A-4683195, US-A-4800195 and US-A-4965188 provides additional uses for oligonucleotides based upon the PDE\_XIV sequence. Such oligomers are generally chemically synthesized, but they may be generated enzymatically or produced from a recombinant source. Oligomers generally comprise two nucleotide sequences, one with sense orientation (5'→3') and one with antisense (3'←5') employed under optimised conditions for identification of a specific gene or condition. The same two oligomers, nested sets of oligomers, or even a degenerate pool of oligomers may be employed under less stringent conditions for detection and/or quantification of closely related DNA or RNA sequences.

The nucleic acid sequence for PDE\_XIV can also be used to generate hybridisation probes as previously described, for mapping the endogenous genomic sequence. The sequence may be mapped to a particular chromosome or to a specific region of the chromosome using well known techniques. These include *in situ* hybridisation to chromosomal spreads (Verma *et al* (1988) Human Chromosomes: A Manual of Basic Techniques, Pergamon Press, New York City), flow-sorted chromosomal preparations, or artificial chromosome constructions such as YACs, bacterial artificial chromosomes (BACs), bacterial PI constructions or single chromosome cDNA libraries.

*In situ* hybridisation of chromosomal preparations and physical mapping techniques such as linkage analysis using established chromosomal markers are invaluable in extending genetic maps. Examples of genetic maps can be found in Science (1995; 270:410f and 1994; 265:1981f). Often the placement of a gene on the chromosome of another mammalian species may reveal associated markers even if the number or arm of a particular human chromosome is not known. New sequences can be assigned to chromosomal arms, or parts thereof, by physical mapping. This provides valuable information to investigators searching for disease genes using positional cloning or other gene discovery techniques. Once a disease or syndrome, such as ataxia telangiectasia (AT), has been crudely localised by genetic linkage to a particular genomic region, for example, AT to 11q22-23 (Gatti *et al* (1988) Nature 336:577-580), any sequences mapping to that



area may represent associated or regulatory genes for further investigation. The nucleotide sequence of the subject invention may also be used to detect differences in the chromosomal location due to translocation, inversion, etc between normal, carrier or affected individuals.

5

## PHARMACEUTICALS

The present invention also provides a pharmaceutical composition for treating an individual in need of same due to PDE\_XIV activity, the composition comprising a therapeutically effective amount of an agent that affects (such as inhibits) said activity and a pharmaceutically acceptable carrier, diluent, excipient or adjuvant.

Thus, the present invention also covers pharmaceutical compositions comprising the agents of the present invention (an agent capable of modulating the expression pattern of the nucleotide sequence of the present invention or the activity of the expression product thereof and/or an agent identified by an assay according to the present invention). In this regard, and in particular for human therapy, even though the agents of the present invention can be administered alone, they will generally be administered in admixture with a pharmaceutical carrier, excipient or diluent selected with regard to the intended route of administration and standard pharmaceutical practice.

By way of example, in the pharmaceutical compositions of the present invention, the agents of the present invention may be admixed with any suitable binder(s), lubricant(s), suspending agent(s), coating agent(s), or solubilising agent(s).

25

In general, a therapeutically effective daily oral or intravenous dose of the agents of the present invention is likely to range from 0.01 to 50 mg/kg body weight of the subject to be treated, preferably 0.1 to 20 mg/kg. The agents of the present invention may also be administered by intravenous infusion, at a dose which is likely to range from 0.001-10 mg/kg/hr.

30

Tablets or capsules of the agents may be administered singly or two or more at a time, as appropriate. It is also possible to administer the agents of the present invention in sustained release formulations.

- 5 Thus, the present invention also provides a method of treating an individual in need of same due to PDE<sub>XIV</sub> activity comprising administering to said individual an effective amount of the pharmaceutical composition of the present invention.

Typically, the physician will determine the actual dosage which will be most suitable for an individual patient and it will vary with the age, weight and response of the particular patient. The above dosages are exemplary of the average case. There can, of course, be individual instances where higher or lower dosage ranges are merited, and such are within the scope of this invention.

- 15 Where appropriate, the pharmaceutical compositions can be administered by inhalation, in the form of a suppository or pessary, topically in the form of a lotion, solution, cream, ointment or dusting powder, by use of a skin patch, orally in the form of tablets containing excipients such as starch or lactose, or in capsules or ovules either alone or in admixture with excipients, or in the form of elixirs, solutions or suspensions containing flavouring or colouring agents, or they can be injected parenterally, for example intracavernosally, intravenously, intramuscularly or subcutaneously. For parenteral administration, the compositions may be best used in the form of a sterile aqueous solution which may contain other substances, for example enough salts or monosaccharides to make the solution isotonic with blood. For buccal or sublingual administration the compositions may be administered in the form of tablets or lozenges which can be formulated in a conventional manner.

For some applications, preferably the compositions are administered orally in the form of tablets containing excipients such as starch or lactose, or in capsules or ovules either alone or in admixture with excipients, or in the form of elixirs, solutions or suspensions containing flavouring or colouring agents.

For parenteral administration, the compositions are best used in the form of a sterile aqueous solution which may contain other substances, for example enough salts or monosaccharides to make the solution isotonic with blood.

- 5 For buccal or sublingual administration the compositions may be administered in the form of tablets or lozenges which can be formulated in a conventional manner.

For oral, parenteral, buccal and sublingual administration to subjects (such as patients), the daily dosage level of the agents of the present invention may typically be from 10 to 500  
10 mg (in single or divided doses). Thus, and by way of example, tablets or capsules may contain from 5 to 100 mg of active agent for administration singly, or two or more at a time, as appropriate. As indicated above, the physician will determine the actual dosage which will be most suitable for an individual patient and it will vary with the age, weight and response of the particular patient. It is to be noted that whilst the above-mentioned  
15 dosages are exemplary of the average case there can, of course, be individual instances where higher or lower dosage ranges are merited and such dose ranges are within the scope of this invention.

In some applications, generally, in humans, oral administration of the agents of the present  
20 invention is the preferred route, being the most convenient and can in some cases avoid disadvantages associated with other routes of administration - such as those associated with intracavernosal (i.c.) administration. In circumstances where the recipient suffers from a swallowing disorder or from impairment of drug absorption after oral administration, the drug may be administered parenterally, e.g. sublingually or buccally.

25

For veterinary use, the agent of the present invention is typically administered as a suitably acceptable formulation in accordance with normal veterinary practice and the veterinary surgeon will determine the dosing regimen and route of administration which will be most appropriate for a particular animal. However, as with human treatment, it may be possible  
30 to administer the agent alone for veterinary treatments.

Typically, the pharmaceutical compositions - which may be for human or animal usage - will comprise any one or more of a pharmaceutically acceptable diluent, carrier, excipient

or adjuvant. The choice of pharmaceutical carrier, excipient or diluent can be selected with regard to the intended route of administration and standard pharmaceutical practice. As indicated above, the pharmaceutical compositions may comprise as - or in addition to - the carrier, excipient or diluent any suitable binder(s), lubricant(s), suspending agent(s),  
5 coating agent(s), solubilising agent(s).

In some embodiments of the present invention, the pharmaceutical compositions will comprise one or more of: an agent that has been screened by an assay of the present invention; an agent that is capable of interacting with any one of the sequences shown in  
10 the attached sequence listings including derivatives, fragments, homologues or variants thereof or sequences capable of hybridising to any one of the sequences shown in the attached sequence listings.

Included in the scope of the invention are oligonucleotide sequences, antisense RNA and  
15 DNA molecules and ribozymes, which function to destabilise PDE\_XIV mRNA or inhibit translation of a PDE\_XIV. Such nucleotide sequences may be used in conditions where it would be preferable to increase cyclic nucleotide levels, such as in inflammation.

A PDE\_XIV antisense molecule may provide the basis for treatment of various abnormal  
20 conditions related to, for example, increased PDE\_XIV activity.

A PDE\_XIV nucleic acid antisense molecule may be used to block the activity of the PDE\_XIV in conditions where it would be preferable to elevate cyclic nucleotide levels.

25 Expression vectors derived from retroviruses, adenovirus, herpes or vaccinia viruses, or from various bacterial plasmids, may be used for delivery of recombinant PDE\_XIV sense or antisense molecules to the targeted cell population. Methods which are well known to those skilled in the art can be used to construct recombinant vectors containing PDE\_XIV. Alternatively, recombinant PDE\_XIV can be delivered to target cells in liposomes.

30

The full length cDNA sequence and/or its regulatory elements enable researchers to use PDE\_XIV as a tool in sense (Youssofian H and HF Lodish 1993 Mol Cell Biol 13:98-104) or antisense (Eguchi *et al* (1991) Annu Rev Biochem 60:631-652) investigations of

gene function. Oligonucleotides, designed from the cDNA or control sequences obtained from the genomic DNA can be used *in vitro* or *in vivo* to inhibit expression. Such technology is now well known in the art, and sense or antisense oligonucleotides or larger fragments can be designed from various locations along the coding or control regions.

5 Appropriate oligonucleotides, which can be 20 nucleotides in length, may be used to isolate PDE\_XIV sequences or closely related molecules from human libraries.

Additionally, PDE\_XIV expression can be modulated by transfecting a cell or tissue with expression vectors which express high levels of a PDE\_XIV fragment in conditions where

10 it would be preferable to block phosphodiesterase activity thereby increasing cyclic nucleotide levels. Such constructs can flood cells with untranslatable sense or antisense sequences. Even in the absence of integration into the DNA, such vectors may continue to transcribe RNA molecules until all copies of the vector are disabled by endogenous nucleases. Such transient expression may last for a month or more with a non-replicating

15 vector and even longer if appropriate replication elements are part of the vector system.

Modifications of gene expression can be obtained by designing antisense sequences to the control regions of the PDE gene, such as the promoters, enhancers, and introns.

20 Oligonucleotides derived from the transcription initiation site, e.g., between -10 and +10 regions of the leader sequence, are preferred. Antisense RNA and DNA molecules may also be designed to block translation of mRNA by preventing the transcript from binding to ribosomes. Similarly, inhibition can be achieved using Hogeboom base-pairing methodology, also known as "triple helix" base pairing. Triple helix pairing compromises

25 the ability of the double helix to open sufficiently for the binding of polymerases, transcription factors, or regulatory molecules.

Thus the invention provides a pharmaceutical composition comprising an agent of the present invention (or even a pharmaceutically acceptable salt thereof, or a pharmaceutically

30 acceptable solvate thereof) together with a pharmaceutically acceptable diluent, excipient or carrier.

The pharmaceutical composition could be for veterinary (i.e. animal) usage or for human usage.

Thus, the present invention therefore also relates to pharmaceutical compositions comprising effective amounts of inhibitors or antagonists of PDE\_XIV protein (including anti-sense nucleic acid sequences) in admixture with a pharmaceutically acceptable diluent, carrier, excipient or adjuvant (including combinations thereof).

The present invention relates to pharmaceutical compositions which may comprise all or portions of PDE\_XIV polynucleotide sequences, PDE\_XIV antisense molecules, PDE\_XIV polypeptides, protein, peptide or organic modulators of PDE\_XIV bioactivity, such as inhibitors, antagonists (including antibodies) or agonists, alone or in combination with at least one other agent, such as stabilising compound, and may be administered in any sterile, biocompatible pharmaceutical carrier, including, but not limited to, saline, buffered saline, dextrose, and water.

#### GENERAL METHODOLOGY REFERENCES

Although in general the techniques mentioned herein are well known in the art, reference may be made in particular to Sambrook *et al.*, Molecular Cloning, A Laboratory Manual (1989) and Ausubel *et al.*, Short Protocols in Molecular Biology (1999) 4<sup>th</sup> Ed, John Wiley & Sons, Inc. PCR is described in US-A-4683195, US-A-4800195 and US-A-4965188.

DEPOSITS

5 The following samples were deposited in accordance with the Budapest Treaty at the recognised depositary The National Collections of Industrial and Marine Bacteria Limited (NCIMB) at 23 St. Machar Drive, Aberdeen, Scotland, United Kingdom, AB2 1RY on 18 December 1998:

*E. coli* pHS-PDE\_XIV NCIMB number NCIMB 40995

*E. coli* pMM-PDE\_XIV NCIMB number NCIMB 40996

10

The following sample was deposited in accordance with the Budapest Treaty at the recognised depositary The National Collections of Industrial and Marine Bacteria Limited (NCIMB) at 23 St. Machar Drive, Aberdeen, Scotland, United Kingdom, AB2 1RY on 9 September 1999:

15

NCIMB number NCIMB 41027 is *E. coli* pHS-PDE\_XIVhm

20

The present invention also encompasses sequences derivable and/or expressable from those deposits and embodiments comprising the same. The present invention also encompasses partial sequences derivable and/or expressable from those deposits and embodiments comprising the same, wherein those partial sequences code for active enzymatic sites. The present invention also encompasses proteins comprising sequences derivable and/or expressable from those deposits and embodiments comprising the same. The present invention also encompasses proteins comprising partial sequences derivable and/or

25 expressable from those deposits and embodiments comprising the same, wherein those partial sequences code for active enzymatic sites.

The present invention also encompasses sequences derivable and/or expressable from those deposits and embodiments comprising the same.

00445-12345-67890

## INTRODUCTION TO THE EXAMPLES SECTION AND THE FIGURES

The present invention will now be described, by way of example only, with reference to the accompanying drawings in which:-

5

Figure 1 which presents a photographic image

Figure 2 which presents a photographic image;

10 

Figure 3 which presents a Table and a photographic image;

Figure 4 which presents an alignment of nucleotide sequences (here the human sequence is a truncated sequence);

15 

Figure 5 which presents an alignment of protein sequences (here the human sequence is a truncated sequence); and

Figure 6 which presents a graph.

20 **EXAMPLES****Northern Hybridisation and Probe preparation**

Northern blots, obtained from Clontech, were prehybridised for 1 hour in ExpressHyb  
25 hybridisation solution (Clontech) at 55°C before a radiolabelled PDE\_XIV fragment (DNA  
was labelled using the Megaprime random labelling system (Amersham) strictly following  
the manufacturers instructions with 50uCi of <sup>32</sup>P-dATP) was added to fresh Expresshyb  
and hybridised to the blot overnight at 55°C, with gentle shaking. Blots were then washed  
3X at room temperature for 10 minutes each in 2XSSC (150 mM NaCl, 30 mM Na.citrate)  
30 followed by 2 washes in 0.2XSSC (15 mM NaCl, 3 mM Na.citrate) at 55°C for 20 minutes  
each. Blots were then exposed to autoradiographic film.



**Polymerase Chain Reactions (PCR)**

PCRs were performed using standard reagents and conditions. Briefly, all reaction buffers and enzymes were obtained in kit format from either Clontech (for rapid amplification of cDNA ends (RACE) reactions) or from Life technologies for standard PCR. Oligonucleotides were obtained from a commercial supplier (OSWEL DNA services) and used at a concentration of 400nM. Reactions were performed on a MJ Research PTC-200 thermal cycler, using cycling parameters as recommended by the manufacturer of the kit being used.

**The identification of PDE\_XIV.**

The clone (IMAGE clone id 1364394) was obtained from Research Genetics (2130, Memorial Pkwy, SW Huntsville, AL 358801, USA) as a stab culture in L-agar. Bacteria from the stab culture were streaked onto a 37 mm L-agar plate in the presence of ampicillin at a concentration of 100mg/ml. After overnight growth at 37°C a single clone was picked, using sterile technique, into 5ml of LB-broth containing ampicillin at a concentration of 100 mg/ml and grown at 37°C overnight with shaking (220 rpm). Plasmid DNA was isolated from the bacteria using a standard commercially available miniprep kit (Qiagen™) and strictly following the manufacturers instructions. The isolated DNA was then subjected to full length sequencing, using standard proprietary kits and reagents and an ABI (PE Applied Biosystems Incorporated) automated sequencer.

**Isolation of a putative full length cDNA clone for PDE\_XIV**

To facilitate the isolation of a murine full length cDNA for PDE\_XIV containing the full coding region and a terminator codon, the expression of PDE\_XIV was determined in a range of tissues using Northern hybridisation. These data (Figures 1 and 2) show that whilst the messenger RNA (length 5.5kb), which hybridises to PDE\_XIV, is present in several tissues, it is particularly highly expressed in murine brain, skeletal muscle & liver and at the following stages of embryological development, 10.5, 13.5, 15.5 days. The 13.5 day murine embryonic cDNA library was used in a cDNA positive selection method (GeneTrapper, Life Technologies, Inc.) to isolate the PDE\_XIV. The method was carried

out as manufacturers instructions. The original EST sequence of clone 1364394 was in the reverse orientation, therefore, design of the selection oligonucleotide needed to be identical to the coding strand running 5' to 3', this step was critical to the success of the cloning strategy. The oligonucleotide used for the selection and repair is detailed below.

5

Oligo 1:

Select1 5'-GGT CAC AGA ACT GCC ACT ATG GTT AAA TGT- 3'

- 10 To isolate the human homologue of the murine PDE\_XIV, PCR primers were designed based on the known EST 1364394 sequence, (Primer 1 and Primer 2) and used to screen a panel of cDNA libraries. PCR's carried out using these primers (the PCR was set up using reagents obtained from a PCR Reagent System kit (Life Technologies) and following the manufacturers instructions, with standard cycling parameters) resulted in the generation of
- 15 a fragment of the expected size, 164bp, obtained from the human HELA cell line cDNA library(Life Technologies, Inc.), using the same selection/repair oligonucleotide (Oligo 1) the human clone was isolated.

Primer 1:

20

newPDE1 5' -ACC GCT CAG AGA TTT CAC AGC A - 3'

Primer 2:

25

newPDE2 5' -CCC GTC TGA CCC CTT AGT CGT A -3'

- Clones were selected for follow up through screening by colony PCR using the above primers and the following method. Using a sterile toothpick a single colony was picked and added to a tube containing 25ul of 1\*PCR supermix (Life Technologies, Inc) & 200nM
- 30 of primers. The tube was vortex'ed for 1 second, centrifuged for 2 seconds at 14,000 x g. The tubes were then placed in a pre-warmed thermal cycler (MJ Research PTC-200 thermal cycler) 94°C. PCR was performed using the following program: 1 cycle; 94°C, 1

minute, followed by 30 cycles of: 94°C, 30 seconds, 55°C, 30 seconds, 72°C, 1 minute. PCR products were then analysed by gel electrophoresis using standard methods.

5 Analysis of the isolated murine clone allowed the assembly of a contiguous sequence of 2823 bp which contains an ORF of 446 residues. The full length cDNA sequence for murine PDE\_XIV is presented as SEQ ID No. 7 with the translation of the largest open reading frame within this cDNA presented as SEQ ID No. 1. The coding region is presented as SEQ ID No. 2.

10 Analysis of the isolated human clone allowed the assembly of a contiguous sequence of 2992 bp which contains an ORF of 288 residues. The cDNA sequence for human PDE\_XIV is presented as SEQ ID No. 8 with the translation of the largest open reading frame within this cDNA presented as SEQ ID No. 3. The coding region is presented as SEQ ID No. 4. This clone has been used to screen a master RNA blot (Clontech) using  
15 standard Northern blot methods to identify the tissue distribution for this cDNA.

The results (see Figure 3) show that the transcript is highly expressed in the putamen and Caudate nucleus of the brain, in addition to Occipital lobe of the brain, heart, ovary, pituitary gland, kidney, liver, small intestine, thymus, and appendix.

20 Like all mammalian phosphodiesterases sequenced to date PDE\_XIV contains a conserved catalytic domain sequence of approximately 250 amino acids in the carboxyl-terminal half of the protein that is thought to be essential for catalytic activity. This segment comprises amino acids 108 to 413 in SEQ ID 1 and exhibits sequence conservation with the  
25 corresponding region of other PDE's. From the nucleotide alignment (see Figure 4) and the protein alignment (see Figure 5) it is clear to see that the human PDE\_XIV clone is truncated at the 3' end due to a premature stop codon at position 1102bp.

#### **Isolation of a putative full length human cDNA clone for PDE\_XIV**

30 A human foetal brain cDNA library was purchased from Life Technologies, Inc (LTI). 100ml of Luria Broth (10g tryptone, 5g yeast extract, 5g NaCl/liter) + 100 ug/ml Ampicillin was inoculated with  $2.5 \times 10^9$  c.f.u of each cDNA library. Each library was

grown up and prepared individually. The cultures were grown up overnight at 30°C and plasmid DNA was prepared as described in the GeneTrapper™ protocol (LTI). Detailed methods for the positive selection of cDNA clones using GeneTrapper™ (LTI) are outlined in the kit protocols, these were followed except for the following modifications. The single stranded cDNA library was combined with 20ng of biotinylated capture oligonucleotide and hybridised at 37°C for 1 hour. The eluted single stranded DNA was repaired using the capture oligonucleotide as a primer for DNA polymerase extension. The repaired cDNA libraries were electroporated in to the *E.coli* strain DH10B (LTI) using the manufacturers protocol.

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### **Cloning, tissue distribution and sequence analysis of human PDE\_XIV**

To facilitate the isolation of a full length cDNA, the expression profile of PDE\_XIV was determined in a range of mouse tissues using Northern hybridisation. This identified a transcript of 5.0 kb which is particularly highly expressed in the mouse embryo at 13.5 days and in the heart, brain and liver of the adult mouse. This transcript is also present at low levels in 8.5, 10.5, 15.5 day embryos and in skeletal muscle, kidney and lung of the adult. A transcript was not detected in the spleen or testis. Based on the data a 13.5 day embryo cDNA library (LTI) was used as a template to isolate the full length sequence using a positive cDNA selection technique, GeneTrapper (LTI). This was achieved by using a PDE\_XIV specific oligonucleotide. A PCR was carried out and we were able to identify cDNAs which contained at least the EST fragment. Restriction endonuclease digestion identified a number of clones containing an insert of 2885bp and full sequence analysis showed that these clones encoded an ORF of 446 amino acids with a predicted molecular mass of 51.3 kDA.

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In addition, the full length human cDNA was isolated by GeneTrapper using the same specific capture oligonucleotide and a human foetal brain cDNA library as template. A number of clones were isolated containing an insert of 2653bp and full sequence analysis identified an ORF of 450 residues with a predicted molecular mass of 51.8 kDA. Using the full length human cDNA an RNA master blot was probed. This identified that the mRNA for human PDE\_XIV is highly expressed in the caudate nucleus, putamen and occipital lobe of the brain and peripherally in the heart, ovary, pituitary gland, kidney, liver, small

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intestine and thymus. Lower levels were observed in skeletal muscle, colon, bladder, uterus, prostate, stomach, adrenal gland, thyroid gland. The transcript was not observed in testis, spleen, pancreas, peripheral leukocytes, lymph node, bone marrow, aorta or cerebellum.

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Sequence comparison of the mouse and human PDE\_XIV shows that they share 91% amino acid identity. The predicted human ORF is extended by 4 amino acids, this is the result of an insertion of five amino acids at position 428 and the deletion of a single amino acid at position 441 in the mouse sequence. Furthermore, both cDNAs contain a single cAMP protein kinase phosphorylation site at serine 45. A phylogenetic alignment of the 230 amino acid catalytic domain of PDE\_XIV (amino acids 172-420) with representatives of other PDEs shows that PDE\_XIV has highest homology to and clusters with PDE7A (about 70% identity). Further bioinformatic analysis shows that this domain shares less than 50% sequence identity with the equivalent regions in the other nine known PDE sub-families and thus may indicate that PDE\_XIV may be a second representative of the PDE7 family.

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The PDE\_XIV sequence contains two putative divalent cation binding motifs HXXXXH<sub>8</sub><sub>20</sub>D (amino acids 173-180, 213-270), which are believed to be essential for hydrolytic activity. Hence, this is further evidence that PDE\_XIV encodes a functional phosphodiesterase enzyme.

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#### **Phosphodiesterase assays on crude mammalian cell lysates**

Both the murine & human PDE\_XIV cDNA clones were transfected into mammalian COS7 cell line (ATCC) using Lipofectamine (Life Technologies, Inc) and standard methods. The COS7 cells were harvested 72 hr's post transfection and assayed for PDE activity using the following phosphodiesterase activity, a commercially available SPA (scintillation proximity assay) kit (Amersham) for cAMP SPA assay (Amersham). Serial dilution's of the cell lysates were made in lysis buffer (50 mM HEPES pH 7.2, 1.92 mM MgCl<sub>2</sub>, 50 mM KCL, 10 mM EGTA, 1 protease inhibitor tablet/50mls) to dilute out any endogenous inhibitory effects. To 25 ml of diluted crude lysate, 25 ml of buffer D (Buffer C + BSA @ 2mg/ml) was added followed by 50 ml of buffer C (20 mM Tris.HCL (pH

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7.4), 5 mM  $\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$ ) containing 500 nM of cAMP substrate, (20  $\mu\text{l}$  of [ $^3\text{H}$ ] cAMP (1  $\mu\text{M}$ , 4  $\mu\text{Ci/ml}$ ) plus 423  $\mu\text{l}$  of cold cAMP, 10  $\mu\text{M}/5\text{ml}$ ). The reaction was incubated for 30 minutes at 30°C. 50  $\mu\text{l}$  of SPA beads was then added, immediately centrifuged at 2000 rpm for 5 minutes. Reading were collected on a Topcount (Wallac) scintillation counter.

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The results (see Figure 6) illustrate that both the human and murine PDE\_XIV clones to be active, 10 fold higher than the mock transfected control.

### **Phosphodiesterase expression in Baculovirus**

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PDE\_XIV was expressed in Baculovirus when fused to an N-terminal FLAG-tag - details on which are presented later.

### **Phosphodiesterase cAMP activity**

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Using cAMP Scintillation Proximity Assays on purified expressed PDE\_XIV with cAMP concentrations ranging from 0 - 10 $\mu\text{M}$ , the expressed PDE\_XIV exhibited a strong cAMP activity with a  $K_m$  of 0.08 $\mu\text{M}$  (as determined by a Hanes plot derived from the kinetic curve). Studies also revealed that the PDE\_XIV has no activity against cGMP. Thus, the

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### **Phosphodiesterase inhibition studies**

PDE\_XIV was found not to be sensitive to the PDE inhibitors milrinone, Rolipram and Ariflo. However, inhibition of PDE\_XIV activity was seen with Dipyridamole and IBMX with  $\text{IC}_{50}$  values of 10.72 $\mu\text{M}$  and 9.32 $\mu\text{M}$  respectively. The data for these studies are presented in the following Table.

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EFFECTS OF INHIBITORS ON PDEXIV		
COMPOUND	SELECTIVITY	IC <sub>50</sub> FOR PDEXIV (μM) (N=2)
Dipyridamole	PDE5/6/8 (0.9/0.38/1.5μM)	10.72 ± 2.74
IBMX	Non-selective (2-50μM)	9.72 ± 0.32
Rolipram	PDE4 (2.0μM)	>30
Milrinone	PDE3/4 (3.2μM/19μM)	>30
Ariflo	PDE4D (40nM)	>30

### Baculovirus expression

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The following studies demonstrate that the PDE enzyme of the present invention - called PDE here for short - can be generated using a baculovirus expression system.

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The following studies also demonstrate that the PDE shows cyclic nucleotide hydrolytic activity when it has been expressed in the baculovirus system.

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The PDE enzyme was generated using the baculovirus expression system based on *Autographa californica* nuclear polyhedrosis virus (AcNPV) infection of *Spodoptera frugiperda* insect cells (*Sf9* cells).

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In these studies, cDNA encoding PDE was cloned into the donor plasmid pFASTBAC-FLAG which contains a mini-Tn7 transposition element. The recombinant plasmid was transformed into DH10BAC competent cells which contain the parent bacmid bMON14272 (AcNPV infectious DNA) and a helper plasmid. The mini-Tn7 element on the pFASTBAC donor can transpose to the attTn7 attachment site on the bacmid thus introducing the PDE gene into the viral genome. Colonies containing recombinant bacmids are identified by disruption of the *lacZ* gene. The PDE/bacmid construct can then be isolated and infected into insect cells (*Sf9* cells) resulting in the production of infectious recombinant baculovirus particles and expression of recombinant PDE-FLAG fusion protein.

25

The phosphodiesterase activity of the crude cell extracts was measured.

Cells were harvested and extracts prepared 24, 48 and 72 hours after transfection.

These results confirm that PDE cDNA encodes a phosphodiesterase which is able to  
5 hydrolyse cAMP.

The crude lysate material was purified by FPLC using a column containing agarose beads  
(M2 affinity gel) to which a purified IgG<sub>1</sub> monoclonal anti-FLAG antibody had been  
conjugated by hydrazide linkage (Eastman Kodak). This allows the specific retention on  
10 the column of the recombinant material (since this is fused to the FLAG epitope) whilst the  
endogenous insect proteins are washed off in the eluate. The recombinant material is then  
washed off under conditions of low pH. This purified material was more suitable for  
detailed enzymatic and inhibitor studies. The purity of the material is assessed by  
coomassie staining after sodium dodecyl sulphate-polyacrylamide gel electrophoresis  
15 (SDS-PAGE) or through western blotting onto a nitro-cellulose membrane of an unstained  
SDS-PAGE (containing recombinant PDE) and analysis with the IgG1 monoclonal anti-  
FLAG epitope antibody. The PDE-FLAG fusion protein is detected due to the interaction  
between the anti-FLAG antibody and the FLAG epitope which is fused to the PDE  
protein.

20 The phosphodiesterase activity of the purified PDE-FLAG fusion protein was assayed  
using a commercially available SPA (scintillation proximity assay) kit (Amersham -  
Amersham place, Little Chalfont, Bucks, HP7 9NA UK) for either cAMP and/or cGMP  
hydrolytic activity. This can be used to permit the determination of the Km value for PDE  
25 against cAMP by determining the enzyme activity at a range of substrate concentrations  
allowing the calculation of an approximate Vmax value for the enzyme.

The results of these experiments show that the PDE cDNA encodes a phosphodiesterase  
which is able to hydrolyse cAMP.

30



In summary the present invention provides and the Examples show *inter alia*:

- 5 1. Novel amino acids.
2. Novel nucleotide sequences.
3. Assays using said novel sequences.
4. Compounds/compositions identified by use of said assays.
5. Expression systems comprising or expressing said novel sequences.
- 10 6. Methods of treatment based on said novel sequences.
7. Pharmaceutical compositions based on said novel sequences.

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All publications mentioned in the above specification are herein incorporated by reference. Various modifications and variations of the described methods and system of the present invention will be apparent to those skilled in the art without departing from the scope and spirit of the present invention. Although the present invention has been described in  
20 connection with specific preferred embodiments, it should be understood that the invention as claimed should not be unduly limited to such specific embodiments. Indeed, various modifications of the described modes for carrying out the invention which are obvious to those skilled in biochemistry and biotechnology or related fields are intended to be within the scope of the following claims.

25

**CLAIMS**

1. An amino acid sequence comprising the sequence presented as Formula I or a variant,  
5 homologue, fragment or derivative thereof.
2. An amino acid sequence comprising the sequence presented as Formula I.
3. A nucleotide sequence encoding the amino acid sequence as defined in claim 1 or  
10 claim 2.
4. A nucleotide sequence comprising the sequence presented as SEQ ID No. 2 or SEQ  
ID No. 4 or SEQ ID No. 6 or a variant, homologue, fragment or derivative thereof.
- 15 5. A nucleotide sequence comprising the sequence presented as SEQ ID No. 2 or SEQ  
ID No. 4 or SED ID No. 6.
6. A nucleotide sequence that is capable of hybridising to the nucleotide sequence  
according to any one of claims 3 to 5.  
20
7. A nucleotide sequence that is capable of hybridising to the nucleotide sequence  
according to claim 6.
8. A vector comprising the nucleotide sequence according to any one of claims 3 to 7.  
25
9. A host cell into which has been incorporated the nucleotide sequence according to any  
one of claims 3 to 7.
10. An assay method for identifying an agent that can affect PDE<sub>XIV</sub> activity or  
30 expression, the assay method comprising  
  
contacting an agent with an amino acid according to claim 1 or claim 2 or a  
nucleotide sequence according to any one of claims 3 to 7; and

measuring the activity or expression of PDE<sub>XIV</sub>;

wherein a difference between a) PDE activity or expression in the absence of the agent and b)  
5 PDE activity or expression in the presence of the agent is indicative that the agent can affect  
PDE<sub>XIV</sub> activity or expression.

11. An assay method according to claim 10 wherein the assay is to screen for agents  
useful in the treatment of disorders found in any one or more of the putamen, the Caudate  
10 nucleus of the brain, the Occipital lobe of the brain, the heart, ovary, the pituitary gland,  
kidney, liver, small intestine, thymus, skeletal muscle, leukocyte regions, dorsal root  
ganglia, uterus, cochlea, small intestine (duodenum), astrocytoma, and appendix.

12. A process comprising the steps of:  
15

- (a) performing the assay according to claim 10 or claim 11;
- (b) identifying one or more agents that do affect PDE<sub>XIV</sub> activity or expression; and
- 20 (c) preparing a quantity of those one or more identified agents.

13. A method of affecting *in vivo* PDE<sub>XIV</sub> activity or expression with an agent;  
wherein the agent is capable of affecting PDE<sub>XIV</sub> activity or expression in an *in vitro* assay  
25 method;

wherein the *in vitro* assay method is the assay method defined in claim 10 or claim 11.

14. Use of an agent in the preparation of a pharmaceutical composition for the treatment of a  
30 disease or condition associated with PDE<sub>XIV</sub>, the agent is capable of having an effect on the  
activity or expression of PDE when assayed *in vitro* by the assay method according to claim  
10 or claim 11.

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15. An enzyme capable of having an immunological reaction with an antibody raised against PDE\_XIV.
16. A nucleotide sequence coding for a PDE, wherein the nucleotide sequence is obtainable from NCIMB 40995 or NCIMB 40996 or NCIMB 41027.
17. A PDE wherein the PDE is expressable from a nucleotide sequence obtainable from NCIMB 40995 or NCIMB 40996 or NCIMB 41027.
18. Use of an agent which has an effect on the activity of PDE\_XIV or the expression thereof in the preparation of a pharmaceutical composition for the treatment of a disease or condition associated with PDE\_XIV.
19. Use of a PDE\_XIV gene and/or expression product thereof in the preparation of a medicament for the treatment and/or modulation of disturbances associated with an imbalance or disturbance of PDE\_XIV.
20. Use according to claim 19 wherein the PDE\_XIV and/or expression product thereof is used to screen for agents that can modulate the activity of the PDE\_XIV and/or expression product thereof.
21. A PDE\_XIV agonist wherein the PDE\_XIV is as defined in claim 1 or is the nucleotide sequence coding for same.
22. A PDE\_XIV antagonist wherein the PDE\_XIV is as defined in claim 1 or is the nucleotide sequence coding for same.
23. A recombinant PDE\_XIV enzyme.
24. A recombinant nucleotide sequence encoding a PDE\_XIV enzyme.
25. A PDE\_XIV enzyme substantially as described herein.

**ABSTRACT****PHOSPHODIESTERASE ENZYMES**

5

Amino acid sequences and nucleotide sequences relating to PDE\_XIV are described. In a preferred aspect, the amino acid sequences comprise the sequence presented as SEQ ID No. 1 or SEQ ID No. 3 or SEQ ID No. 5.

10

Sequence "B547.4460

Fig 1

Murine MTN Blot  
probed with  $^{32}\text{P}$ -labelled  
murine PDE\_XIV

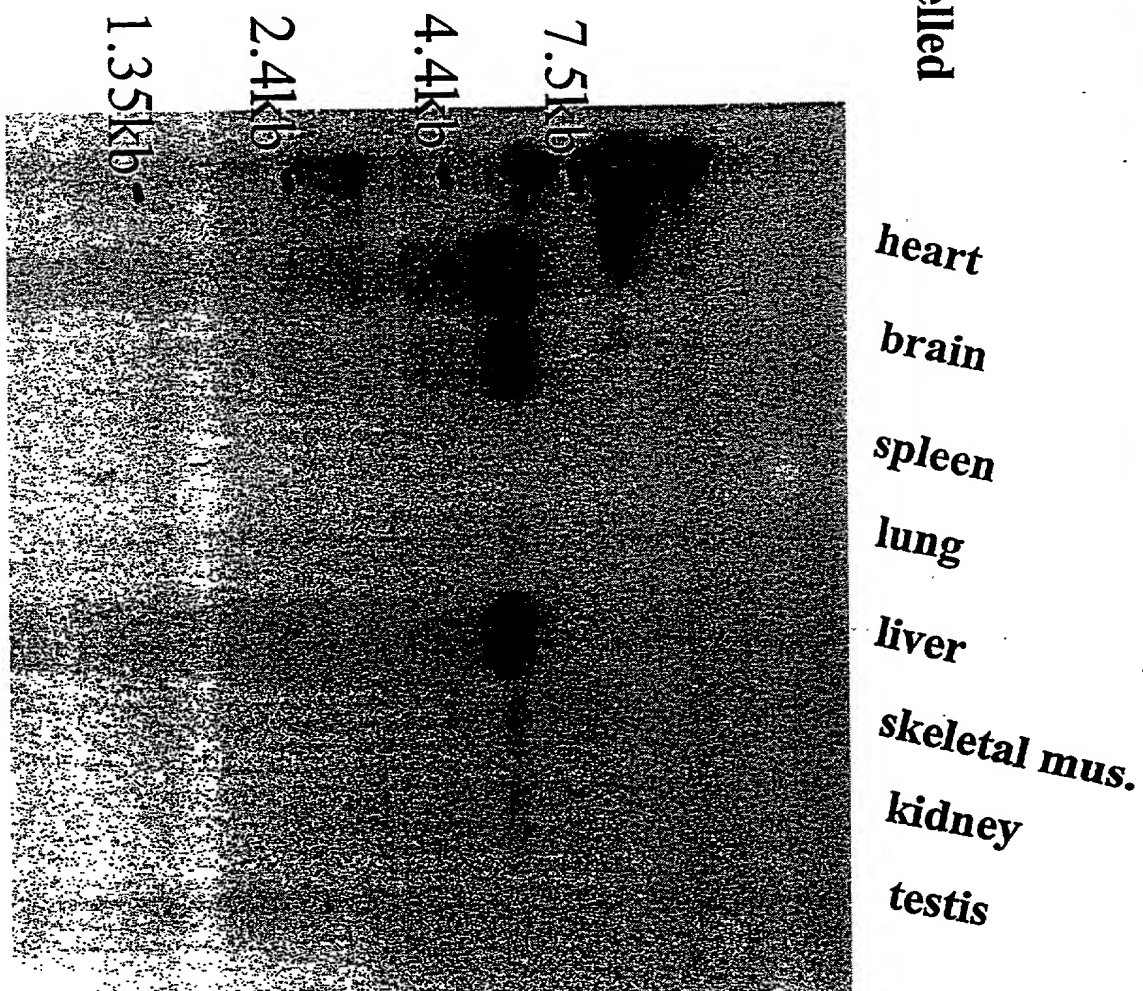
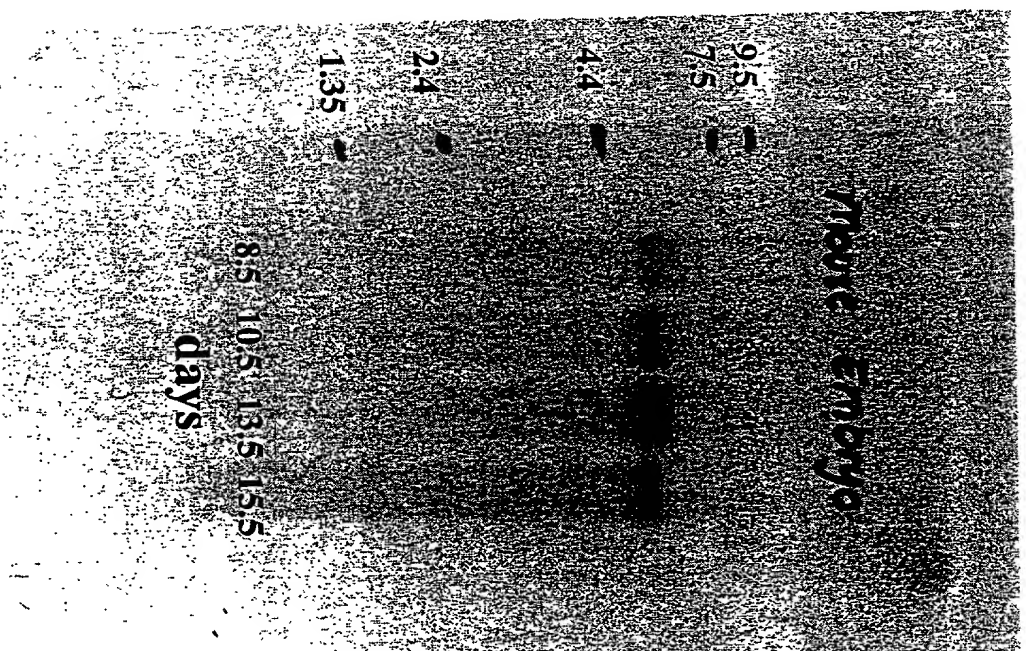


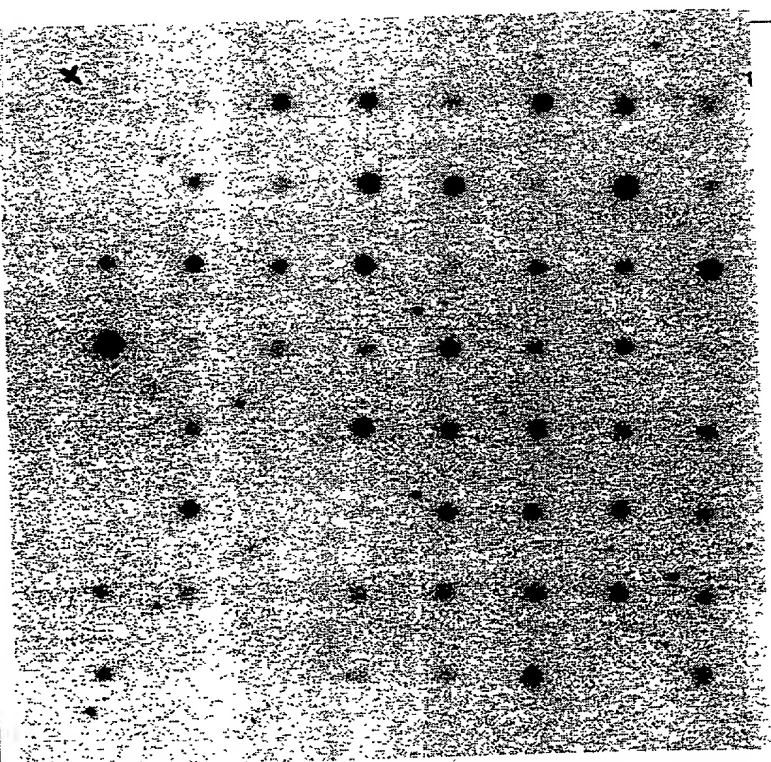
Fig 2

Murine Embryo MTN Blot  
probed with  $^{32}\text{P}$ -labelled  
murine PDE\_XIV



**Figure 3**  
**Human RNA Master Blot probed with  $^{32}\text{P}$ -labelled**  
**human PDE\_XIV**

	1	2	3	4	5	6	7	8
A	whole brain	amygdala	caudate nucleus	cerebellum	cerebral cortex	frontal lobe	hippocampus	medulla oblongata
B	occipital lobe	putamen	substantia nigra	temporal lobe	thalamus	nucleus accumbens	spinal cord	
C	heart	aorta	skeletal muscle	colon	bladder	uterus	prostate	stomach
D	testis	ovary	pancreas	pituitary gland	adrenal gland	thyroid gland	salivary gland	mammary gland
E	kidney	liver	small intestine	spleen	thymus	peripheral leukocyte	lymph node	bone marrow
F	appendix	lung	trachea	placenta				
G	fetal brain	fetal heart	fetal kidney	fetal liver	fetal spleen	fetal thymus	fetal lung	
H	yeast total RNA 100 ng	yeast tRNA 100 ng	<i>E. coli</i> rRNA 100 ng	<i>E. coli</i> DNA 100 ng	Poly (dA) 100 ng	human C <sub>1</sub> 1 DNA 100 ng	human DNA 100 ng	human DNA 500 ng





**Fig. 4. Alignment of the Murine and Human PDE\_XIV nucleotide sequence**

5 New sequence is PDE\_XIV

Pileup: Genetics Computer Group.

MSF: 3134 Type: N Check: 5422 ..

10 Name: mpdea\_oo Len: 3134 Check: 5084 Weight: 0.001  
Name: hspdea\_oo Len: 3134 Check: 338 Weight: 0.100

	new mpde_	1	AGGTACGCCT	GCAGGTACCG	GTCCGGAATT	CCCGGGTCTGA	CCCACGCGTC
	new hspde_		.....	.....	.....	.....	.....
15	new mpde_	51	CGGCCAGCCT	CCCAGGCCGG	CTGCCTGCTC	ACCCAGCCAG	TCGCTAGCTC
	new hspde_	1	.....	.....	.....	.....	.CGGAAT.TC
	mpdea_	101	TGGGCACTGC	AGCAGGCTCG	GCTCTGTCCC	AGCGCTCGCT	TGCTTGCTCG
20	hspde_	9	GATGCACTGC	AGCAGGCTCG	GCTCTGTCCC	AGC....A..	.....
	mpdea_	151	CTCGCTCGGC	TGGGAGAAAA	GTGGTGTC.C	TCGCCCAG..	AGAGCCTCTC
	hspde_	43	....CTTGTC	TGGGAGAAAA	GTGGTGTTAC	TCACCCAGGG	AGAGTCTCTC
25	mpdea_	198	TCTC..CCTT	CCTTCTTTCT	CGAGCTCTCT	GAGTCCTTTG	GCGTTTCTTT
	hspde_	89	TTTCTACCTT	CCTTCTTTCT	CGATCTCCTT	GTGTGCTTTT	GTGTTTCTTT
	mpdea_	246	CTTCTTTTCC	TTTTTTTTTT	TTTTTTAATA	TTTTCTTTTT	CTTCTATAA
30	hspde_	139	ATTTCTTTTC	CTTTTTTTTC	TT.....	TTTTTTTTTT	GTTACT....
	mpde_	296	AACTTGCATA	ATTATACTGC	TAATCCTGGA	TGAGGTTGCT	GGATTCTGCA
	hspde_	177	.....TA	ATTATATTCC	TAATCCTGGA	TGAAGTTGCT	GGATTCTGCA
35	mpde_	346	GCACAAATCT	TCATGAACAA	GCCGCACCGC	TCAGAGATTT	CACAGCATTC
	hspde_	219	GCACAAGTCT	TCATGAACAA	GCAGCACCGC	TCAGAGATTT	CACGGCATTC
					start codon		
	mpde_	396	AAAGGTCACA	GAAGTGCCAC	TATGGTTAA <u>A</u>	TGTCTTGTTT	AATGGTTGAG
40	hspde_	269	AAAGGTCACA	GAAGTGCCAC	TATGGTTAA <u>A</u>	TGTCTTGTTT	AATGGTTGAG
	mpdea_	446	AGGTGTGGCG	AAGTCTTGTT	TGAGAGCCCT	GAACAGAGTG	TCAAATGTGT
	hspde_	319	AGGTGTGGCG	AAATCTTGTT	TGAGAACCCC	GATCAGAATG	CCAAATGTGT
45	mpde_	496	TTGCATGCTA	GGAGATGTAC	GAATAAGGGG	TCAGACGGGG	GTTCTTGCCG
	hspde_	369	TTGCATGCTG	GGAGATATAC	GAATAAGGGG	TCAGACGGGG	GTTCTTGCTG
	mpde_	546	AACGCCGTGG	CTCCTACCCA	TTCATTGACT	TCCGTCTACT	TAACAATACA
	hspde_	419	AACGCCGTGG	CTCCTACCCA	TTCATTGACT	TCCGCCTACT	TAACAGTACA
50	mpde_	596	ACACACTCAG	GGGAAATTGG	CACCAAGAAA	AAGGTGAAAC	GACTGTTAAG
	hspde_	469	ACATACTCAG	GGGAGATTGG	CACCAAGAAA	AAGGTGAAAA	GACTATTAAG
	mpde_	646	TTTCCAAAGA	TACTTCCATG	CATCTAGGCT	TCTCCGGGGG	ATTATACCGC
55	hspde_	519	CTTTCAAAGA	TACTTCCATG	CATCAAGGCT	GCTTCGTGGA	ATTATACCAC
	mpde_	696	AGGCCCCCTCT	CCACCTGCTG	GATGAAGACT	ACCTTGGAACA	AGCAAGGCAC
	hspde_	569	AAGCCCCCTCT	GCACCTGCTG	GATGAAGACT	ACCTTGGAACA	AGCAAGGCAT
60	mpde_	746	ATGCTCTCCA	AAGTTGGAAC	GTGGGACTTT	GACATTTTCT	TGTTTGATCG
	hspde_	619	ATGCTCTCCA	AAGTGGGAAT	GTGGGATTTT	GACATTTTCT	TGTTTGATCG

65222T "65442460

## PC10315AGPR

	mpde_	796	CTTGACAAAT	GGGAACAGTC	TGGTAACTCT	GTTGTGTCAC	CTCTTCAACT
	hspde_	669	CTTGACAAAT	GGAAACAGCC	TGGTAACTACT	GTTGTGCCAC	CTCTTCAATA
5	mpde_	846	CCCATGGGCT	CATCCACCAT	TTCAAGCTCG	ATATGGTGAC	CTTGCACAGG
	hspde_	719	CCCATGGACT	CATTACCAT	TTCAAGTTAG	ATATGGTGAC	CTTACACCGA
	mpde_	896	TTTCTGGTTA	TGGTTCAGGA	AGATTACCAC	GGTCACAACC	CATACCACAA
10	hspde_	769	TTTTTAGTCA	TGGTTCAGA	AGATTACCAC	AGCCAAAACC	CGTATCACAA
	mpde_	946	TGCTGTTTAC	GCAGCCGACG	TCACCCAGGC	CATGCAC TGT	TACCTGAAGG
	hspde_	819	TGCTGTTTAC	GCAGCCGACG	TCACCCAGGC	CATGCAC TGC	TACCTGAAAG
	mpde_	996	AGCCAAAGTT	GGCAAGCTTC	CTCACACCTC	TGGACATCAT	GCTTGGACTA
15	hspde_	869	AGCCAAAGCT	TGCCAGCTTC	CTCACGCCTC	TGGACATCAT	GCTTGGACTG
	mpde_	1046	CTGGCTGCAG	CAGCTCATGA	CGTGGACCAC	CCAGGGGTCA	ACCAGCCATT
	hspde_	919	CTGGCTGCAG	CAGCACACGA	TGTGGACCAC	CCAGGGGTGA	ACCAGCCATT
20	mpde_	1096	TTTGATCAAA	ACTAACCACC	ATCTTGCCAA	CCTGTATCAG	AATATGTCTG
	hspde_	969	TTTGATAAAA	ACTAACCACC	ATCTTGCCAA	CCTATATCAG	AATATGTCTG
	mpde_	1146	TACTGGAGAA	TCACCACTGG	CGATCTACAA	TTGGCATGCT	TCGAGAATCA
25	hspde_	1019	TGCTGGAGAA	TCATCACTGG	CGATCTACAA	TTGGCATGCT	TCGAGAATCA
	mpde_	1196	CGGCTCCTGG	CTCACTTGCC	AAAGGAAATG	ACACAGG...	...ATATC
	hspde_	1069	AGGCTTCTTG	CTCATTTGCC	AAAGGAAATG	ACGTAAGTGC	TGCCGAGATG
						stop codon	
30	mpde_	1238	GAACA.....	....GCA...	.....GCTG	GGCTCCCTCA	TCTTGGCCAC
	hspde_	1119	AAACATACTG	ATGTGCATGC	AGTAAAGATA	AGCCACTTTC	TCTAGGGCA.
	mpde_	1270	GGATATCAAC	AGACAGAATG	AGTTTCTGA.	.....CCCG	CTTAAAAGCT
35	hspde_	1168	GGCTTGGGAC	CTTTTGCGTG	AATGGCAGAG	AGCCCCCGG	CTGTACTTCC
	mpde_	1313	CACCTCCACA	ATAAAGATT.	TGAGAC....	..TGGAGAAT	GT.ACAGGA.
	hspde_	1218	TGCCTGCACT	GAGCTGTCTA	TCAGAGGAGA	TTTGGTGTCA	GTTACAGCAA
	mpde_	1354	..CAGACACT	TTATGCT.TC	AGATCGCCTT	GAAGTGTGCT	GACATTTGCA
40	hspde_	1268	CCCAGAAACC	AAAATCTCTC	TGTGTGCTTT	GAAAGGGCCT	TGCAGAGTCA
	mpde_	1401	AT..CCTT..	GTC.GTATCT	GGGAGATGAG	CAAGCAGT..	GGAGTGAAAG
	hspde_	1318	ATGACCTACA	GTCAGGAAAA	GGGATAATAA	ACAGCTCTCA	GTTTTCACAC
45	mpde_	1444	GGT.....	CTGTGAGGAA	TTCTACAGAC	AAGGTGACCT	TGAAC..AG.
	hspde_	1368	GCTTCAGTAT	CAGTGCTCAA	CTTTGCCAAA	TTCCCGACCT	TTAGTTTAGC
	mpde_	1484	AAGTTTGAAC	TGGAATTCAG	.TCCTCTTTG	TAATCAAC.A	GAAAGATTCA
50	hspde_	1418	AAAATTGTCC	TTCCATGTAG	CTCCAAATAG	TAAATATTTA	TCAAGAAGGA
	mpde_	1532	ATCCCTAGCA	TACAAA...T	TGGTTTCATG	ACT.TACATC	GTGGAGCCGC
	hspde_	1468	A.CCCAGGCA	TTCTAAAGCT	AGAGTTCAAA	AAAGTATATT	TTGTAATTGC
55	mpde_	1578	TGTTCCGGG.	...AGTGG..	GCCCCGTTTA	CTGGG..AAC	AGCACCTGT
	hspde_	1517	TAGTCTCAGC	AAAAATAGAA	GTCAGAAATT	CTTTTCTAAA	ATGTCTTTTG
	mpde_	1620	CGGAGAACAT	GCTAAGCC..	....ATCTCG	CGCACAACAA	AGCCAGTGG
	hspde_	1567	CTAAGTAATT	GAAATGGCCC	TAGCATTTTT	TTACCAATT	AATTTACCTT
60	mpde_	1664	AAGAGCCT.G	CTGTCCAATC	AGCAC...AG	ACGCA.....	...GGGGCAG
	hspde_	1617	ACGTCTCTTG	CACCTTAAAC	AGAAGGGGAG	ACACTCATTT	TCTGGTTTAC
	mpde_	1702	CG.....G	CCAGGACCTG	GCGG....GC	CCCGC...AC	CTGAGACCCT
65	hspde_	1667	TATTTGATAG	CCATGGTATG	TAGGCTGAGT	CCCACTAAAT	CTGAGGCCAT
					stop codon		
	mpde_	1738	GGAG.CAGAC	AGAAGGTGCC	ACGCCCCTAAG	GTAGCTGTC.	.TGCTGA..T
70	hspde_	1717	TGTTTCATTT	TCCTGGTG..	..GCCCCAAG	TTAGCTGCTA	ATACTGTCTT

PC10315AGPR							
	mpde_	1783	GCACGGCCA.	.....TCT	G.TCCGTCCA	.....CAGGA	GCACGGCC..
	hspde_	1763	CCAAGGCCAC	CATTAAATCT	GATCTGTTTA	ATGAACACGT	GCAGAACCCA
5	mpde_	1817	...ATCC...	.....G	TCC...GACT	GC.....	.CCTCGCAAC
	hspde_	1813	AGAAACCTAG	GTGAAAAGAG	TACATAGATT	GCTGTACCCCT	TCTTCAAGAC
	mpde_	1840	AAGCCCATCA	CGCTGGGTTT	CGATGCCAT.	.CCGCCTGCC	A.CTTACC..
	hspde_	1863	AAGCACATAA	CTTGAGGTCA	AGGACCAAGT	GCTGTCTCCC	AACTGAACAA
10	mpde_	1885	...GCCTCCC	TTCGTTGATC	CAAGTGTACA	AAAGCCATTG	...TCACCTC
	hspde_	1913	GCAGTATACT	CTGGGTGTG	GATTGATTCC	TGGCCCTCTG	ATTTGATCTC
	mpde_	1929	AGCAT.....	.TAGCTGCC.	..GAAATGGG	CGGCTCTATC	CCGTCATTGG
	hspde_	1963	ATGCTGTTTC	CTAGCACCCA	GAGGAATGTG	AAATTTGCAG	GAGGAATTTT
15	mpde_	1970	AG..CTGAT.	...TCTGGGG	CGGCTGCCCC	AACCGAAAC.	.....G
	hspde_	2013	AGTTCTGATA	AATTTTACT	CCCTGGAAC	AAATAAAACC	AGTTCTCGTG
	mpde_	2004	CCTGGAAGTA	AGAA..AGGG	GTGCTTCTGC	CGTGTTCGCC	TCTGGCCCTT
	hspde_	2063	CATGGAATAA	AACTTATGC	CTCTTACTAG	AATAATAAAT	TGCAAAAGATT
	mpde_	2052	GGTCACGCTG	ACTGGCAGTA	GCTCCTAAGT	CCAGAGCATT	TTAACGTTTG
	hspde_	2113	GAAAGAATTA	AATGCAAAAA	GAATAAAAA	CTAGAGCAAA	AGATCAAGTG
25	mpde_	2102	CCATC..GGA	CAGCTGACCT	...GCATGAC	ACCAGCAT..	.ACTTGAAC
	hspde_	2163	AGAAGAAGAA	AAGAGGAGGT	AAGGAGAGAG	ACAAGGAAGA	AAGAAGGAGA
	mpde_	2144	TGCAAAACTG	GTCTTGCGTG	CCAGAGCACA	AACGAGAGTG	TGAGAGAAA.
	hspde_	2213	AGGAAAGGAA	GAATAGTCAG	G.ACAGGAAA	GAAGAAAATG	CAAGGGAAAT
30	mpde_	2193	..GTACCTTC	TATTT..TAA	TAATAATTAT	TATTATAAAA	TA....ATAA
	hspde_	2262	GGGAAAGGAC	TCGGGGTGA	CCAGACTTCT	CCTGGTCAGT	ACCTGCATTTC
	mpde_	2235	ATCTTTTTTAA	CTTTT..ATA	TTTCATGCAC	CAGACAATGG	GTCTAAAACT
	hspde_	2312	ATCCTGTTTG	TTACTCAATA	TTTCTTTTCT	AAAATATTCA	TTTCACATCT
	mpde_	2283	TTGGA...CA	AGTAATACTC	TGCGTACCCA	AACCTAAGAG	G.....GGG
	hspde_	2362	ATGGATTCCA	ATGAAAAATA	TATTTTTATG	TGTCTTTGTG	GAACACAGTG
40	mpde_	2324	TTC...ATTA	TTTT.GCTAT	T.GACTC...	..TATGCCAC	ATTGGGTCCG
	hspde_	2412	TTATAAATTG	TTTTTGCCAG	AAGAATAATT	GTATACAAT	AATATATGTG
	mpde_	2364	AGA..TGTGG	CACCATGCG	ATTTCTGAAA	CCACGCGTCC	.CCTCCCATC
	hspde_	2462	AAAACCTTTAT	TACAAAAGCC	ATTATCATAA	TCATTATTAT	TCCTTCTATC
45	mpde_	2411	TGGTGAAGG	TGCTGTACAG	CCCGTCCC..	..TTTGCACC	GTTAGCCAAT
	hspde_	2512	ACA.GGTAAA	TGCTTTAATG	TCATTTTTCT	GATTTTAAAA	GTAGGGCAGG
	mpde_	2457	CCGTCTTT..	....TACGGA	...TTCAGTG	ACCTGTTTAT	ATTACAA.G
	hspde_	2561	TTAATTGTAG	AAAGTAAGGA	AAATTCAGGA	AAGTGTTAGT	TTGAACATG
	mpde_	2497	TGTACATTTT	CTGT..AAAT	ACCAAACGCT	ACTGA.....	.....TTCC
	hspde_	2611	TGAAGTTGCT	CTTTTAAAG	GCCAAAAACA	GGAGACTTTT	AGCACTTTCA
55	mpde_	2534	CATGC..CA.	....AAATAC	ACGAGTATTA	TGGGATTGCT	A.....CCTG
	hspde_	2661	TATGTTTCAG	CTTGATATGA	AAGAGAAAAC	TGAAACTGCT	AGTAATCCTG
	mpde_	2571	.....T	ATAAACATG	GCACTGTGAA	CAGAATA...	.CTGTTAGTT
	hspde_	2711	CCATCCAGGT	ATAGTTCATG	TTAACCTGGC	TAGTTTATTT	TCTTTTAGTC
60	mpde_						
	hspde_						

1. *Chlorophyll a* (Chl a) content (mg/g dry weight) was determined using a spectrophotometer (Shimadzu UV-1601) at 663 nm. The absorbance was converted to concentration using a standard curve.

15

Figure 5. Protien Alignment of the Murine &amp; Human PDE\_XIV

CLUSTAL W (1.7) multiple sequence alignment

```

5  mpde      MSCLMVERCGEVLFPESPEQSVKVCMLGDVRLRGQTGVPAERRGSPYFIDFRLLNNTTHS 60
   hspde     MSCLMVERCGEILFENFDQNAKVCMLGDIRLRGQTGVPAERRGSPYFIDFRLLNSTTYS 60
          *****:***.*:*.*****:***** *****:***.*

10  mpde      GEIGTKKKVKRLLSFQRYFHASRLLRGLIPQAPLHLLDEOYLGQARHMLSKVGTWDFDIF 120
   hspde     GEIGTKKKVKRLLSFQRYFHASRLLRGIIPQAPLHLLDEOYLGQARHMLSKVGMWDFDIF 120
          *****:***.*:*.*****:***** *****:***.*

15  mpde      LFDRLTNGNSLVTLTLLCHLFNSHGLIHHFKLDMVTLHRFLVMVQEDYHGHNPYHNAVHAAD 180
   hspde     LFDRLTNGNSLVTLTLLCHLFNSHGLIHHFKLDMVTLHRFLVMVQEDYHSQNPYHNAVHAAD 180
          *****:***.*:*.*****:***** *****:***.*

20  mpde      VTQAMHCYLKEPKLASFLTPLDIMLGLLAAAAHVDVHPGVNQPFLLIKTNHHLANLYQNMS 240
   hspde     VTQAMHCYLKEPKLASFLTPLDIMLGLLAAAAHVDVHPGVNQPFLLIKTNHHLANLYQNMS 240
          *****:***.*:*.*****:***** *****:***.*

25  mpde      VLENHHWRSTIGMLRESRLLAHLPKEMTQDIEQQLGSLILATDINRQNEFLTRLKAHLHN 300
   hspde     VLENHHWRSTIGMLRESRLLAHLPKEMT----- 268
          *****:***.*:*.*****:***** *****:***.*

30  mpde      KDLRLENVQDRHFMQLKALCADICNPCIWEMSKQWSEVCEEFYRQGDLEQKFELEIS 360
   hspde     ----- 320

35  mpde      PLCNQKQKDSIPSIQIGFMTYIVEPLFREWARFTGNSTLSENMLSHLAHNKAQWKSLLENQ 420
   hspde     ----- 380

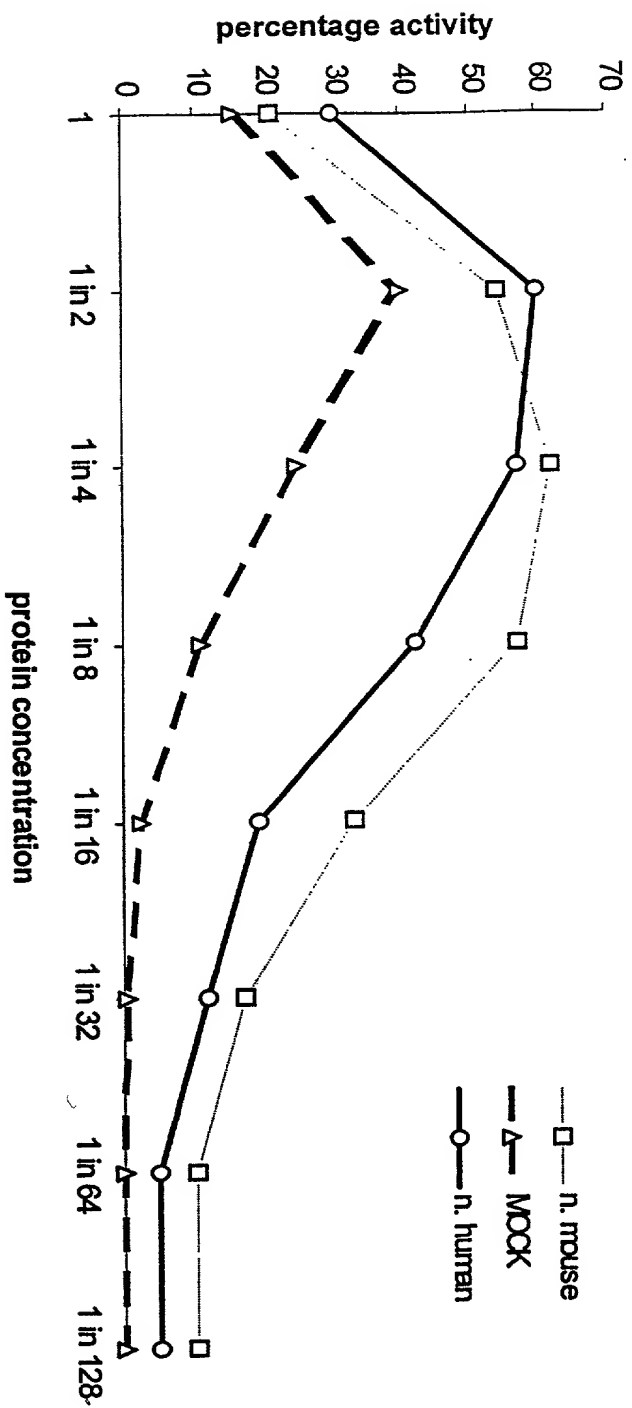
   mpde      HRRRGSGQDLGAPAPETLEQTECATP 446
   hspde     ----- 400

```

Zinc binding motif's are highlighted in bold.  
Non-Catalytic domain in italics.

**Figure 6.**

**SPA assay to determine the cAMP hydrolytic  
activity of murine and  
human PDE\_XIV.**



Please type a plus sign (+) inside this box →

+

<b>DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (37 CFR 1.63)</b>  <input checked="" type="checkbox"/> Declaration submitted with Initial Filing <input type="checkbox"/> Declaration Submitted after Initial Filing (surcharge 37 CFR 1.16 (e) required)	<b>Attorney Docket Number</b>	PC10315AGPR
	<b>First Named Inventor</b>	Mark D. Fidock
	<b>COMPLETE IF KNOWN</b>	
	<b>Application Number</b>	To be assigned.
	<b>Filing Date</b>	Herewith
	<b>Group Art Unit</b>	To be assigned.
	<b>Examiner Name</b>	To be assigned.

**As a below named inventor, I hereby declare that:**

My residence, post office address, and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Enzyme PDE xiv

(Title of the Invention)

the specification of which

☒ is attached hereto

OR

☐ was filed on (MM/DD/YYYY) \_\_\_\_\_ as United States Application Number or PCT International

Application Number \_\_\_\_\_ and was amended on (MM/DD/YYYY) \_\_\_\_\_ (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or of any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached?	
				YES	NO
9828603.2	Great Britain	December 23, 1998	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
9922123.6	Great Britain	September 17, 1999	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

☐ Additional foreign application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto:

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**U.S. Parent Application Number or PCT Parent Number**

**Parent Filing Date (MM/DD/YYYY)**

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James T. Jones	30,561	A. Dean Olson	31,185
Gregg C. Benson	30,997	Mervin E. Brokke	32,723
Robert F. Sheyka	31,304	Valerie M. Fedowich	33,688
Grover F. Fuller Jr.	31,760	Bryan C. Zielinski	34,462
Karen DeBenedictis	32,977	Robert T. Ronau	36,257
Lorraine B. Ling	35,251	B. Timothy Creagan	39,156
Garth Butterfield	36,997	Alan L. Koller	37,371
Carl J. Goddard	39,203	Jolene W. Appleman	35,428
Raymond M. Speer	26,810	Kristina L. Konstas	37,864
Jennifer A. Kispert	40,049	Gregory P. Raymer	36,647
Martha A. Gammill	31,820	Jacob M. Levine	32,509
Israel Nissenbaum	27,582	Seth H. Jacobs	32,140
Steven W. Collier	42,429	E. Victor Donahue	35,492
Roy F. Waldron	42,208	Todd M. Crissey	37,807

☐ Additional registered practitioner(s) named on supplemental Registered Practitioner Information sheet PTO/SB/02C attached hereto.

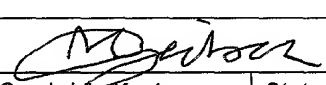
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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

**Name of Sole or First Inventor:** ☐ A petition has been filed for this unsigned inventor

<b>Given Name (first and middle [if any])</b>			<b>Family Name or Surname</b>				
Mark D.			Fidock				
<b>Inventor's Signature</b>					<b>Date</b>	16 December 1999	
<b>Residence: City</b>	Sandwich, Kent	<b>State</b>		<b>Country</b>	England	<b>Citizenship</b>	Great Britain
<b>Post Office Address</b>	Ramsgate Road, Sandwich, Kent, CT13 9NJ						
<b>Post Office Address</b>							
<b>City</b>	Sandwich, Kent	<b>State</b>		<b>Zip</b>	CT13 9NJ	<b>Country</b>	England

☐ Additional inventors are being named on the attached a supplemental Additional Inventor(s) sheet(s) PTO/SB/02A attached hereto.



SEQ ID No. 1 = murine amino acid sequence

SEQ ID No. 3 = truncated human amino acid sequence

SEQ ID No. 5 = human amino acid sequence

15

SEQ ID No. 8 = truncated human cDNA sequence

(It is to be noted that in the above text, references to SEQ ID No. 2 are equally applicable to SEQ ID No. 7. Also, references to SEQ ID No. 4 or SEQ ID No. 6 are equally applicable to SEQ ID No. 8.)

MSCLMVERCGEVLFESEPEQSVKVCMLGDVRLRGQTVPAERRGSYPFIDFRLNNTTHSGEIGTKKKVK 70  
RLLSFQRYFHASRLLRGIIPQAPLHLLDEDEDYLGQARHMLSKVGTWDFDIFLFDRLTNGNSLVTLCHLNF 140  
5 SHGLIHHFKLDMVTLHRLVLMVQEDYHGHNPNYHNAVHAADVTOAMHCYLKEPKLASFLTPLDIMLGLLAA 210  
AAHDVDHDPGVNQPFLLIKTNHHLNLYQNMSVLENNHWRSTIGMLRESRLLAHLPKEMTQDIEQQGLSLIL 280  
ATDINRQNEFLTRLKAHLHNKDLRLENVQDRHFMLQIALKCADICNPCIWEMSKQWSERVEEYFQRQGD 350  
LEQKFELEISPLCNQQKDSIPSIQIGFMTYIVEPLFREWARFTGNSTLSENMLSHLAHNKAQWKSLLSNQ 420  
HRRRSGSQDLGAPAPETLEQTEGATP 446

SEQ ID No. 2

start  
15 ATGTCTTGTGTTTAATGGTTGAGAGGTGTGGCGAAGTCTTGTGTTGAGAGCCCTGAACAGAGTGTCAAA  
TGTGTTTGCATGCTAGGAGATGTACGACTAAGGGGTCAGACGGGGGTTCTGCCGAACGCCGTGGCTCCT  
ACCCATTTCATTGACTTCCGTCTACTTAACAATACAACACACTCAGGGGAAATGGCACCAAGAAAAAGGT  
GAAACGACTGTAAAGTTTCCAAAGATACCTCCATGCATCTAGGCTTCTCCGGGGGATTATACCGCAGGCC  
CCTCTCCACCTGCTGGATGAAGACTACCTTGGACAAGCAAGGCACATGCTCTCCAAAGTTGGAACGTGGG  
ACTTTGACATTTTCTTGTGTTGATCGCTTGACAAATGGGAACAGTCTGGTAACCTCTGTTGTGCACCTCTT  
20 CAACTCCCATGGGCTCATCCACCATTTCAGCTCGATATGGTGACCTTGACAGGTTTCTGGTTATGGTT  
CAGGAAGATTACCACGGTCACAACCCATACCACAATGCTGTTACGCAGCCGACGTCACCCAGGCCATGC  
ACTGTTACCTGAAGGAGCCAAAGTTGGCAAGCTTCTCACACCTCTGGACATCATGCTTGACTACTGGC  
TGCAGCAGCTCATGACGTGGACCACCCAGGGGTCAACCAGCCATTTTGTGATCAAACTAACCACCATCTT  
GCCAACCTGTATCAGAATATGCTGTACTGGAGAATCACCCTGGCGATCTACAATTGGCATGCTTCGAG  
25 AATCACGGCTCCTGGCTCACTTGCCAAAGGAAATGACACAGGATATCGAACAGCAGCTGGGCTCCCTCAT  
CTTGGCCACGGATATCAACAGACAGAATGAGTTTCTGACCCGCTTAAAAGCTCACCTCCACAATAAAGAT  
TTGAGACTGGAGAATGTACAGGACAGACACTTTATGCTTCAGATCGCCTTGAAGTGTGCTGACATTTGCA  
ATCCTTGTCTGTATCTGGGAGATGAGCAAGCAGTGGAGTGAAAGGGTCTGTGAGGAATTCTACAGACAAGG  
TGACCTTGAACAGAAGTTTGAAGTGGAAATCAGTCCCTCTTTGTAATCAACAGAAAGATTCAATCCCTAGC  
30 ATACAAATTGGTTTCATGACTTACATCGTGGAGCCCTGTTCGGGAGTGGGCCCCGTTTACTGGGAACA  
GCACCTGTGCGGAGAACATGCTAAGCCATCTCGGCACACAAAGGCCAGTGGAAAGAGCCTGTGTCCAA  
TCAGCACAGACGACGGGCGAGCGCCAGGACCTGGCGGGCCCCGCACCTGAGACCTGGAGCAGACAGAA  
GGTGCCACGCCCTTAA  
stop

SEQ ID No. 3

MSCLMVERCGEILFENPDQNAKVCMLGDIRLRGQTVRAERRGSYPFIDFRLNNTTYSGEIGTKKKVK 70  
RLLSFQRYFHASRLLRGIIPQAPLHLLDEDEDYLGQARHMLSKVGMWDFDIFLFDRLTNGNSLVTLCHLNF 140  
40 THGLIHHFKLDMVTLHRLVLMVQEDYHSQNPYHNAVHAADVTOAMHCYLKEPKLASFLTPLDIMLGLLAA 210  
AAHDVDHDPGVNQPFLLIKTNHHLNLYQNMSVLENNHWRSTIGMLRESRLLAHLPKEMTGTWDFDIFLFDRL 280  
LTNGNSLV 288

SEQ ID No. 4

start  
50 ATGTCTTGTGTTTAATGGTTGAGAGGTGTGGCGAAATCTTGTGTTGAGAACCCCGA  
TCAGAATGCCAAATGTGTTTGCATGCTGGGAGATATACGACTAAGGGGTCAGACGGGGGTTCTGTGCTGAA  
CGCCGTGGCTCCTACCCATTTCATTGACTTCCGCCTACTTAACAGTACAACATACTCAGGGGAGATTGGCA  
CCAAGAAAAAGGTGAAAAGACTATTAAGCTTTCAAAGATACTTCCATGCATCAAGGCTGCTTCGTGGAAT  
TATACCACAAGCCCTCTGCACCTGCTGGATGAAGACTACCTTGGACAAGCAAGGCATATGCTCTCCAAA  
GTGGGAATGTGGGATTTTGACATTTTCTTGTGTTGATCGCTTGACAAATGGAAACAGCCTGGTAACACTGT  
55 TGTGCCACCTCTTCAATACCCATGGACTCATTACCATTTTCAAGTTAGATATGGTGACCTTACACCGATT  
TTTAGTCATGGTTCAAGAAGATTACCACAGCCAAAACCCGTATCACAATGCTGTTACGCAGCCGACGTC  
ACCCAGGCCATGCACTGCTACCTGAAAGAGCCAAAGCTTGCCAGCTTCTCACGCCTCTGGACATCATGC  
TTGGACTGCTGGCTGCAGCAGCACACGATGTGGACCACCCAGGGGTGAACCAGCCATTTTGTATAAAAC  
TAACCACCTCTTGCAACCTATATCAGAATATGCTGTGCTGGAGAATCATCACTGGCGATCTACAATT  
60 GGCATGCTTCGAGAATCAAGGCTTCTTGCTCATTTGCCAAAGGAAATGACGTAA  
stop

SEQ ID No. 5 is the amino acid sequence presented as HS\_PDEXIV. For reference the sequence MM\_PDEXIV is SEQ ID No. 1. CLUSTAL W (1.74) multiple sequence alignment was used.

5

10

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```
HS_PDEXIV      MSCLMVERCGEILFENPDQNAKVCVCM LGDIRLRGQTGVRAERRGSYPFIDFRLNSTTYS
MM_PDEXIV      MSCLMVERCGEVLFESEPEQSVKVCVCM LGDVRLRGQTGVPAERRGSYPFIDFRLNNTTHS
*****:***.*:..*****.***** *****.***:*

HS_PDEXIV      GEIGTKKKVKRLLSFQRYFHASRLLRGIIPQAPLHLLDE DYLGQARHMLS KVGCMWDFDIF
MM_PDEXIV      GEIGTKKKVKRLLSFQRYFHASRLLRGIIPQAPLHLLDE DYLGQARHMLS KVGCTWDFDIF
*****.*****.*****.*****.***** *****

HS_PDEXIV      LFDRLTNGNSLVTL LCHLFNTHGLIHHFKLDMVTLHRFLVMVQEDYHSONPYHNAVHAAD
MM_PDEXIV      LFDRLTNGNSLVTL LCHLFNTHGLIHHFKLDMVTLHRFLVMVQEDYHGHNPYHNAVHAAD
*****.*****.*****.*****.***** *****

HS_PDEXIV      VTQAMHCYLKEPKLASFLTPLDIMLGLLAAA AHDVDHPGVNQPF LIKTNNHHLANLYQNMS
MM_PDEXIV      VTQAMHCYLKEPKLASFLTPLDIMLGLLAAA AHDVDHPGVNQPF LIKTNNHHLANLYQNMS
*****.*****.*****.*****.***** *****

HS_PDEXIV      VLENHHWRSTIGMLRESRLLAHLPKEMTQDIEQQLGSLILATDINRQNEFLTRLKAHLHN
MM_PDEXIV      VLENHHWRSTIGMLRESRLLAHLPKEMTQDIEQQLGSLILATDINRQNEFLTRLKAHLHN
*****.*****.*****.*****.***** *****

HS_PDEXIV      KDLRLEDAQDRHFMLQIALKCADICNPCR IWEMSKQWSERVCEEFYRQGELEQKFELEIS
MM_PDEXIV      KDLRLENVQDRHFMLQIALKCADICNPCR IWEMSKQWSERVCEEFYRQGDLEQKFELEIS
*****.:*****.*****.*****.***** *****

HS_PDEXIV      PLCNQKQKSIPSIQIGFMSYIVEPLFREWAHFTGNSTLSENMLGHLAHNKAQWKSLLPRQ
MM_PDEXIV      PLCNQKQKSIPSIQIGFMTYIVEPLFREWARFTGNSTLSENMLSHLAHNKAQWKSLLSNQ
*****.:*****.*****.*****.***** *****

HS_PDEXIV      HRRRGSGSGPDHDHAGQGTESEEQ-EGDSP
MM_PDEXIV      HRRRGSG-----QDLAGPAPETLEQTEGATP
** ***. : * ** ..: ** ** :*
```

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## SEQ ID No. 6

5 SEQ ID No. 6 is the nucleotide sequence presented as HS\_PDEXIV. For reference the sequence MM\_PDEXIV is SEQ ID No. 2. CLUSTAL W (1.74) multiple sequence alignment was used.

```
10  HS_PDEXIV      -----ATGTCTTGTTTAATGGTTGAGAGGTGTGGCGAAATCTTGTTTGAGAACCCC
    MM_PDEXIV      -----ATGTCTTGTTTAATGGTTGAGAGGTGTGGCGAAGTCTTGTTTGAGAGCCCT
                               *****

    HS_PDEXIV      GATCAGAATGCCAAATGTGTTTGCATGCTGGGAGATATACGACTAAGGGGTGACACGGGG
    MM_PDEXIV      GAACAGAGTGTCAAATGTGTTTGCATGCTAGGAGATGTACGACTAAGGGGTGACACGGGG
    15  ** **** * *****

    HS_PDEXIV      GTTCGTGCTGAACGCCGTGGCTCCTACCCATTCATTGACTTCCGCCTACTTAACAGTACA
    MM_PDEXIV      GTTCCTGCCGAACGCCGTGGCTCCTACCCATTCATTGACTTCCGTCTACTTAACAATACA
    20  **** * *****

    HS_PDEXIV      ACATACTCAGGGGAGATTGGCACCAGAAAAAGGTGAAAAGACTATTAAGCTTTCAAAGA
    MM_PDEXIV      ACACACTCAGGGGAAATGGCACCAAGAAAAAGGTGAAACGACTGTAAAGTTCCAAAGA
    **** * *****

    25  HS_PDEXIV      TACTTCCATGCATCAAGGCTGCTTCGTGGAATTATACCACAAGCCCCCTGTCACCTGCTG
    MM_PDEXIV      TACTTCCATGCATCTAGGCTTCTCCGGGGGATTATACCGCAGGCCCTCTCCACCTGCTG
    *****

    HS_PDEXIV      GATGAAGACTACCTTGGACAAGCAAGGCATATGCTCTCCAAAGTGGGAATGTGGGATTTT
    30  MM_PDEXIV      GATGAAGACTACCTTGGACAAGCAAGGCACATGCTCTCCAAAGTGGAACTGGGACTTT
    *****

    HS_PDEXIV      GACATTTTCTGTTTGATCGCTTGACAAATGGAACAGCCTGGTAACACTGTGTGTGCCAC
    35  MM_PDEXIV      GACATTTTCTGTTTGATCGCTTGACAAATGGGAACAGTCTGGTAACCTGTGTGTGCAC
    *****

    HS_PDEXIV      CTCTTCAATACCCATGGACTCATTCACCATTTCAGTATAGATATGGTGACCTTACACCGA
    40  MM_PDEXIV      CTCTTCAACTCCCATGGGCTCATCCACCATTTCAGCTCGATATGGTGACCTTGCACAGG
    *****

    HS_PDEXIV      TTTTGTAGTCATGGTTCAAGAAGATTACCACAGCCAAAACCCGTATCACAATGCTGTTTAC
    45  MM_PDEXIV      TTTCTGGTTATGGTTTCAAGAAGATTACCACGGTCAACAACCCATACCACAATGCTGTTTAC
    *** * *****

    HS_PDEXIV      GCAGCCGACGTCACCCAGGCCATGCAC'TGCTACCTGAAAGAGCCAAAGCTTGCCAGCTTC
    50  MM_PDEXIV      GCAGCCGACGTCACCCAGGCCATGCAC'TGTTACCTGAAGGAGCCAAAGTTGGCAAGCTTC
    *****

    HS_PDEXIV      CTCACGCCTCTGGACATCATGCTTGGA'CTGCTGGCTGCAGCAGCACACGATGTGGACCAC
    55  MM_PDEXIV      CTCACACCTCTGGACATCATGCTTGGA'CTACTGGCTGCAGCAGCTCATGACGTGGACCAC
    *****

    HS_PDEXIV      CCAGGGGTGAACCAGCCATTTTGTATAAAACTAACCACCATCTTGCAAACCTATATCAG
    60  MM_PDEXIV      CCAGGGGTCAACCAGCCATTTTGTATCAAAACTAACCACCATCTTGCAAACCTGTATCAG
    *****

    HS_PDEXIV      AATATGTCTGTGCTGGAGAATCATCACTGGCGATCTACAATTGGCATGCTTCGAGAATCA
    65  MM_PDEXIV      AATATGTCTGTACTGGAGAATCACCCTGGCGATCTACAATTGGCATGCTTCGAGAATCA
    *****

    HS_PDEXIV      AGGCTTCTTGCTCATTTGCCAAAGGAAATGACACAGGATATTGAACAGCAGCTGGGCTCC
    70  MM_PDEXIV      CGGCTCCTGGCTCACTTGCCAAAGGAAATGACACAGGATATCGAACAGCAGCTGGGCTCC
    **** * *****

    HS_PDEXIV      TTGATCTTGGAACAGACATCAACAGGCAGAATGAATTTTGTACCAGATTGAAAGCTCAC
    75  MM_PDEXIV      CTCATCTTGCCACGGATATCAACAGACAGAATGAGTTCTGACCCGCTTAAAGCTCAC
    * *****

    HS_PDEXIV      CTCCACAATAAAGACTTAAGACTGGAGGATGCACAGGACAGGCACTTTATGCTTCAGATC
    80  MM_PDEXIV      CTCCACAATAAAGATTGAGACTGGAGAAATGTACAGGACAGACACTTTATGCTTCAGATC
    *****

    HS_PDEXIV      GCCTTGAAGTGTGCTGACATTTGCAATCCTTG'TAGAATCTGGGAGATGAGCAAGCAGTGG
    85  MM_PDEXIV      GCCTTGAAGTGTGCTGACATTTGCAATCCTTG'GCGTATCTGGGAGATGAGCAAGCAGTGG
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	CGGAATTCGATGCACCTGCAGCAGGCTCGGCTCTGTCCCAGCACTTGTCTGGGAGAAAAGTGGTGTACTC	70
	ACCCAGGGAGAGTCTCTCTTTCTACCTTCCTTCTCTCGATCTCCTTGTGTGCTTTTGTGTTTCTTTAT	140
	TTCTTTTCTCTTTTCTTTTCTTTTCTTTTGTACTTAATATATCTTAATCTGGATGAAGTTGCTGG	210
	ATTCTGCAGCACAAAGTCTTCATGAACAAGCAGCACCCTCAGAGATTTCACGGCATTCAAAGGTCACAGA	280
5	start	
	ACTGCCACTATGGTTAAATGCTCTTGTAAATGGTTGAGAGGTGTGGCGAAATCTGTTTGAGAACCCCGA	350
	TCAGAATGCCAAATGTGTTTGCATGCTGGGAGATATACGACTAAGGGGTGAGACGGGGGTTCGTGCTGAA	420
	CGCGTGGCTCCTACCCATTTCATTGACTTCCGCTTAAACAGTACAACATACTCAGGGGAGATTGGCA	490
	CCAAGAAAAAGGTGAAAAGACTATTAAAGCTTCAAAGATACTTCCATGCATCAAGGCTGCTTCGTGGAAT	560
10	TATACCACAAGCCCCCTGTCACCTGCTGGATGAAGACTACCTTGGACAAGCAAGGCATATGCTCTCCAAA	630
	GTGGGAATGTGGGATTTTGACATTTCTTGTGTTGATCGCTTGACAAATGGAAACAGCCTGGTAACACTGT	700
	TGTGCCACCTCTTCAATACCCATGGACTCATTCACCATTTCAAGTTAGATATGGTGACCTTACACCGATT	770
	TTTAGTCATGGTTCAAGAAGATTACCACAGCCAAAACCGTATCACAATGCTGTTCACGCAGCCGACGTC	840
	ACCCAGGCCATGCACCTGCTACCTGAAAGAGCCAAAAGCTTGCCAGCTTCCTCAGCCCTCTGGACATCATGC	910
15	TTGGACTGCTGGCTGTCAGCAGCACGATGTGGACCACCCAGGGGTGAACCAGCCATTTTGTATAAAAC	980
	TAACCACCATCTTGCAACCTATATCAGAATATGCTGTGCTGGAGAATCATCACTGGCGATCTACAATT	1050
	stop	
	GGCATGCTTCGAGAATCAAGGCTTCTTGCTCATTTTGCCAAAGGAAATGACGTAAAGTGTGCTGCCGAGATGAA	1120
	ACATACTGATGTGCATGCAGTAAAGATAAGCCACTTCTCTAGGGCAGGCTTGGGACCTTTTGCCTGAAT	1190
20	GGCAGAGAGCCCCCGCTGTACTTCCTGCTGCACCTGAGCTGTCTATCAGAGGAGATTTCGTGTGCTGTT	1260
	ACAGCAACCCAGAAACCAAAATCTCTCTGTGTGCTTTGAAAGGGCCTTGACAGTCAATGACCTACAGTC	1330
	AGGAAAAGGGATAAATAAACAGCTCTCAGTTTTCACACGCTTCAGTATCAGTGTCAACTTTGCCAAAATTC	1400
	CCGACCTTTAGTTTAGCAAAATTTGCTCTCCATGTAGCTCCAAATAGTAAATATTTATCAAGAAGGAACC	1470
	CAGGCATTCTAAAGCTAGAGTTCAAAAAAGTATATTTTGAATTGCTAGTCTCAGCAAAAAATAGAAGTCA	1540
25	GAAATTCCTTTCTAAAAATGTCCTTTGCTAAGTAATTGAAATGGCCCTAGCATTTTTTTCCCAATTAATT	1610
	TACCTTACGTCCTTTGCACTTTAAACAGAGGGGAGACACTCATTTTCTGGTTCATTTTGTATAGCCAT	1680
	GGTATGTAGGCTGAGTCCCACTAAATCTGAGGCCATTGTTTCATTTTCTGGTGGCCCCAAGTTAGCTGC	1750
	TAATACTGTCCTTCCAAAGGCCACCATTAATCTGATCTGTTAATGAACACGTGCAGAACCCAAAGAAACCT	1820
	AGGTGAAAAGAGTACATAGATTGCTGTACCCCTCTTCAAGACAAGCACATAACTTGAGGTCAAGGACCAA	1890
30	GTGCTGTCTCCCACTGAACAAGCAGTATACCTCTGGGTTGTGGATTGATTCTCTGGCCCTCTGATTTGATC	1960
	TCATGCTGTTTCTTAGCACCCAGAGGAATGTGAAATTTGCAGGAGGAATTTTCAGTCTGTATAAATTTTTA	2030
	CTCCCTGGAATAAAATAAACAGTTCTCGTGCATGGAATAAAAACTTATGCCTCTTACTAGAATAATAA	2100
	ATTGCAAAAGATTGAAAGAATTAATATGCAAAAAGAACTAAAACTAGAGCAAAAGATCAAGTGAGAAGAAG	2170
	AAAAGAGGAGCTAAGGAGAGAGACAAGGAAGAAAGAGGAGAAGGAAAGGAAGAATAGTGAGGACAGGAA	2240
35	AGAAGAAAAATGCAAGGGAAATGGGAAAGGACTCTGGGGTGACCAGACTTCTCTGGTCAGTACCTGCATT	2310
	CATCCTGTTTGTACTCAATATTTCTTTCTTAAATATTCATTTTCACATCTATGGATTCCAATGAAAAT	2380
	ATATTTTATGTGCTTTTGTGGAACACAGTGTATAAATTTGTTTGTGCCAGAAGAATAATTGTTATACAA	2450
	TAATATATGTGAAAACCTTTATTACAAAAGCCATTATCATAATCATTATTATTCCTTCTATCACAGGTAAA	2520
	TGCTTTAATGTCATTTTCTGATTTTAAAGTAGGGCAGGTTAATTGTAGAAAGTAAGGAAAATTCAGGA	2590
40	AAGTGTAGTTTGAACATATGTGAAGTTGCTCTTTTAAAGGGCCAAAACAGGAGACTTTTAGCACTTTCA	2660
	TATGTTTCAGCTTGATATGAAAGAGAAAACGAACTGCTAGTAATCCTGCCATCCAGGTATAGTTTCATG	2730
	TTAACCTGGCTAGTTTATTTCTTTTAGTCTTTTTCATAACAACTTATTTTAACAAAATATGATTATA	2800
	TTTGGGGAACCTATTTTACAGTTTACGTCCTGAAATTTTATTATTACAATAAAGACTTTTTCCAAATCA	2870
45	AAAAAAGGGGGCGGCTCTAGAGGATCCCTCGAGGGGCCAAGCTTACGCGTCATGCGACG	2940
	TCATAGCTCTCTCCCTATAGTGAGTCGTATTATAAGCTAGGCACTGGCCGTC	2992

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See Formula I

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